

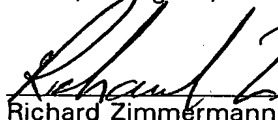
JOINT INVENTORS

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Richard Zimmermann

APPLICATION FOR UNITED STATES LETTERS PATENT

SPECIFICATION

TO ALL WHOM IT MAY CONCERN:

Be it known that we, Lewis B. Schwartz, a citizen of the United States of America, residing at 510 N. Clay Street, Hinsdale 60521-3214, in the County of Cook and State of Illinois; Ralph R. Weichselbaum, a citizen of the United States of America, residing at 1909 N. Burling Street, Chicago 60614-5123, in the County of Cook and State of Illinois; and Bernard Roizman, a citizen of the United States of America, residing at 5555 S. Everett Street, Chicago 60637-1968 in the County of Cook and State of Illinois have invented a new and useful GENETICALLY ENGINEERED HERPES VIRUS FOR THE TREATMENT OF CARDIOVASCULAR DISEASE, of which the following is a specification.

GENETICALLY ENGINEERED HERPES VIRUS FOR THE TREATMENT OF CARDIOVASCULAR DISEASE

The United States Government may own certain rights in this invention.

5

Priority is claimed to U.S. Provisional Patent Application No. 60/253,680,
filed November 28, 2000.

BACKGROUND OF THE INVENTION

10 FIELD OF THE INVENTION

The present invention relates generally to the fields of genetic therapy. More particularly, it concerns administration of a herpes simplex viral vector to produce a therapeutic benefit in vascular and cardiovascular tissue. In additional aspects, the invention concerns combination therapies comprising administration of a herpes simplex viral vector and treatment with at least one additional pharmacological agent or surgical procedure, as well as use of surgical procedures and other targeting means to facilitate delivery of the herpes simplex virus vectors to vascular cells and tissues.

15 RELATED TECHNOLOGY

Vascular disease remains the leading cause of death and disability in the Western world [McGovern *et al.*, *New Engl. J. Med.* **334**:884-890, 1996]. Current treatment strategies are primarily aimed at risk factor modification and/or mechanical remediation of critical lesions. Although these strategies are often effective, the ability to genetically alter the basic pathophysiologic defects within diseased vascular tissue would offer a new paradigm of therapy and possibly revolutionize the treatment of vascular disease. The feasibility of vascular gene transfer was first demonstrated in 1989 when it was shown endothelial cells (EC's) expressing the retrovirally transduced lacZ gene could adhere and function in porcine iliac arteries [Nabel *et al.*, *Science* **244**:1342-1344, 1989]. Since then, nearly one billion U.S. dollars annually have been spent to refine and improve systems of transfer vectors and delivery systems [Svensson *et al.*, *Curr. Opin. Cardio.* **13**:369-374, 1998].

A variety of vectors have been used to transfer genes to vascular tissue, with variable results. Each of the most popular vector systems - naked plasmid DNA [Isner *et al.*, *Hum. Gene Ther.* 7:989-1011, 1996; Baumgartner *et al.*, *Circ.* 97:1114-1123, 1998], liposome-encapsulated DNA, [Armeanu *et al.*, *Mol. Ther.* 1:366-375, 2000] retrovirus [Geary *et al.*, *Hum. Gene Ther.* 5:1211-1216, 1994], adeno-associated virus (AAV) [Lynch *et al.*, *Circ. Res.* 80:497-505, 1997], and adenovirus [Kay, *J. Vasc. Surg.* 24:160-161, 1996; Yeh *et al.*, *FASEB J.* 11:615-623, 1997], have met with some success depending on the application. Efforts at gene transfer using only naked plasmid DNA have met with limited success due to low transfection efficiency [Baumgartner *et al.*, *Circ.* 97:1114-1123, 1998; Reissen *et al.*, *Hum. Gene Ther.* 4:449-458, 1993; Turunen *et al.*, *Gene Ther.* 6:6-11, 1999]. DNA encapsulated within liposomes is somewhat more efficient [Armeanu *et al.*, *Mol. Ther.* 1:366-375, 2000; Turunen *et al.*, *Gene Ther.* 6:6-11, 1999; Takeshita *et al.*, *J. Clin. Invest.* 93:652-661, 1994; Lim *et al.*, *Circ.* 83:2007-2011, 1991; Matsumoto *et al.*, *J. Vasc. Surg.* 27:135-144, 1998; Matsumura *et al.*, *J. Surg. Res.* 85:339-345, 1999] but generally lacks complete penetrance with an intact basement membrane. Retrovirus was tested in early experiments [Nabel *et al.*, *Science* 244:1342-1344, 1989; Dunn *et al.*, *Circ.* 99:3199-3205, 1996], as well as more recently to transfer antisense oligonucleotides [Zhu *et al.*, *Circ.* 96:628-635, 1997], but suffers greatly from its low efficiency in non-dividing cells. A relatively new vector system, adeno-associated virus (AAV), has been shown to efficiently infect skeletal muscle without inciting an intense immune response [Muzyczka, *Curr. Topics Microbiol. Immunol.* 158:97-129, 1992], and preliminary results have demonstrated that AAV is capable of transfecting vascular EC's [Lynch *et al.*, *Circ. Res.* 80:497-505, 1997; Rolling *et al.*, *Gene Ther.* 4:757-761, 1997; Kotin, *Hum. Gene Ther.* 5:793-801, 1994; Xiao *et al.*, *J. Virol.* 70:8098-8108, 1996] smooth muscle cells (SMC's, Rolling *et al.*, *Gene Ther.* 4:757-761, 1997), and cardiocytes [Svensson *et al.*, *Circ.* 99:201-205, 1999]. A recent report, however, indicates that gene transfer into vascular tissue with an intact endothelium may be problematic, with only 1-14% of cells staining positive after 30 days [Eslami *et al.*, *J. Vasc. Surg.* 31:1149-1159, 2000].

Certainly, the most widely tested vector for vascular gene transfer has been replication-deficient adenovirus [Yeh *et al.*, *FASEB J.* 11:615-623, 1997]. Adenoviral vectors exhibit a high penetrance in non-dividing cells, can be produced in high concentration, and can house relatively large transgenes. They have been used experimentally in vascular tissue to transfer a wide variety of marker and biologically active genes including Rb, [Chang *et al.*, *J. Clin. Invest.* 95:2260-2268, 1995; Smith *et al.*, *Circ.* 96:1899-1905, 1997] p21, [Chang *et al.*, *J. Clin. Invest.* 95:2260-2268, 1995; Scheinman *et al.*, *J. Vasc. Surg.* 29:360-369, 1999] cytosine deaminase [Harrell *et al.*, 1997; Fortunato *et al.*, 2000], C-type natriuretic peptide (Ueno *et al.*, 1997), metalloproteinase inhibitors (Cheng *et al.*, 1998; George *et al.*, 2000; Dollery *et al.*, 1999), hirudin, (Bishop *et al.*, 1999) cyclooxygenase (Zoldhelyi *et al.*, 1996), tissue factor pathway inhibitor (Zoldhelyi *et al.*, 2000), thrombomodulin (Waugh *et al.*, 2000), ecNOS (Varenne *et al.*, 1998; Cable *et al.*, 1999), β ARK_{CT}, (Fulton *et al.*, 1996) and antisense RNA to bFGF (Hanna *et al.*, 1997; Neschis *et al.*, 1998; Hanna *et al.*, 2000). The success of many of these protocols has been due, in part, to the very high concentration of vector used, as well as use of the popular model of creating injury by balloon inflation, which denudes endothelium and enhances vector penetration.

Gene transfer into intact vascular tissue, such as non-injured arteries or vein grafts, has proved more difficult (Eslami *et al.*, 2000; Schwartz *et al.*, 1999; Fulton *et al.*, 1996; Hanna *et al.*, 2000; Mann *et al.*, 1995; Fulton *et al.*, 1997; Fulton *et al.*, 1998; Faries *et al.*, 2000). Transfer efficiency is generally lower, the required doses are high, and the biologic effects less pronounced (Schwartz *et al.*, 1999; Fulton *et al.*, 1996; Hanna *et al.*, 2000; Mann *et al.*, 1995; Fulton *et al.*, 1997; Fulton *et al.*, 1998; Faires *et al.*, 2000). Furthermore, and perhaps most importantly, an intact immune system remains a serious impediment to long-term transgene expression using adenovirus, as neutralization of the cytosolic episome generally is complete after the first week (Eslami *et al.*, 2000; Chang *et al.*, 1995). Immunomodulation of the host can extend the life of the transgene [Ascher *et al.*, *Ann. Vasc. Surg.* 14:385-392, 2000], but global immunosuppression in the elderly population with vascular disease is clearly impractical. The problems inherent in adenoviral-based gene transfer in vascular tissue is best illustrated by the fact that,

to date, only a single trial of therapeutic gene transfer has been reported [for angiogenesis, Rosengart et al., 1999], and there has yet to be a published clinical trial using adenoviral vectors for the treatment or prevention of vascular proliferative disorders.

5 A potential vector for the generation of long-term transgene expression is herpes simplex virus type one (HSV-1). HSV-1 is the virus that is responsible for recurrent oropharyngeal cold sores. It is a lytic, nonintegrating DNA virus that has demonstrated neurotropism and the establishment of long-term infections. Like
10 adenovirus, HSV-1 can be manufactured in high titer and contains multiple non-essential genes that can potentially be deleted and replaced with large transgenes. This had led to speculation that the use of HSV may become a method for gene transfer for systemic disease (Culver, 1996; Lachmann and Efstathiou, 1997). However, the lytic and highly infectious nature of HSV, and the lifelong potential for latency, necessitate significant genetic modification prior to therapeutic use
15 (Lachmann and Efstathiou, 1997; Huard et al., 1995; Advani *et al.*, *Gene Therapy* 5:160-165, 1998]. The first set of recombinant herpesviruses to be generated lacked one or more genes (*e.g.*, thymidine kinase or ribonucleotide reductase), which only moderately reduced viral growth in non-dividing cells (Martuza et al., 1991; Mineta et al., 1994; Miyatake et al., 1999). HSV-1 has been extensively studied as therapy
20 for malignant tumors of the central nervous system using a variety of genetic manipulations [Advani *et al.*, *Gene Therapy* 5:160-165, 1998; Chambers et al., 1995; Andreansky *et al.*, *Can. Res.* 57:1502-1509, 1997]. More recently, attempts to render the virus nonvirulent have been made through deletion of both copies of the $\gamma_{134.5}$ gene (Chou et al., 1990; Chou and Roizman, 1994).

25 However, the inventors are only aware of two phase I clinical trials using transfer vectors and delivery systems, both addressing the feasibility of transfer of angiogenic factors [Baumgartner *et al.*, *Circ.* 1114-1123, 1998; Isner et al., 1995; Symes et al., 1999; Rosengart et al., 1999). Thus, despite the advances in the art, there is still a need for more effective vectors and methods of therapy for a wider
30 application of somatic gene therapy to treat vascular disease.

SUMMARY OF THE INVENTION

The present invention provides a solution to the aforementioned problem by providing a method of expressing a heterologous nucleic acid in a vascular cell via the use of a recombinant herpes simplex virus. The present invention also provides
5 a method of treating or preventing vascular disease as well as a method directed to inducing normal physiology in a functionally abnormal vascular

According to one aspect of the invention a method of expressing a heterologous nucleic acid sequence in a vascular cell is provided. The method comprises administering to the vascular cell a recombinant replicating herpes
10 simplex viral vector, wherein the herpes simplex virus comprises a heterologous nucleic acid, and wherein the herpes simplex virus is debilitated for growth in the central nervous system.

The recombinant herpes simplex virus is debilitated for growth via non-silent insertion, substitution, or deletion of a nucleotide sequence in at least one non-
15 essential gene of the herpes simplex virus. In a related embodiment, the recombinant herpes simplex virus may further comprise a non-silent insertion, substitution, or deletion of a nucleotide sequence in at least one essential gene of the herpes simplex virus. In one embodiment, the herpes simplex virus lacks one expressible $\gamma_{134.5}$ gene, a non-essential gene. In a further embodiment, the
20 recombinant herpes simplex lacks both expressible $\gamma_{134.5}$ genes.

Suitable vascular cells for treatment by the present method would include an endothelial cell, a smooth muscle cell, and/or an adventitial cell. In a further embodiment the heterologous nucleic acid sequence encodes a polypeptide, *i.e.*, an antiproliferative polypeptide, a vasodilatory polypeptide, and an angiogenic
25 polypeptide. In an additional embodiment the heterologous nucleic acid sequence encodes an antisense oligonucleotide or antisense polynucleotide. In a further embodiment, the antisense oligonucleotide or antisense polynucleotide is complementary to an RNA encoding an antiproliferative polypeptide, vasodilatory polypeptide, or angiogenic polypeptide. Preferably, the recombinant herpes simplex
30 virus is HSV-1. In a related embodiment, the recombinant herpes simplex virus further comprises at least one gene essential for the treatment of a herpes simplex virus infection by an anti-viral agent.

According to another aspect of the present invention, a method is provided for treating or preventing a cardiovascular disease or condition in a vascular cell comprising administering to the cell a recombinant replicating herpes simplex viral vector comprising a heterologous nucleic acid sequence, and wherein the herpes simplex virus is debilitated for growth in the central nervous system.

The recombinant herpes simplex virus is debilitated for growth via non-silent insertion, substitution, or deletion of a nucleotide sequence in at least one non-essential gene of the herpes simplex virus. The recombinant herpes simplex virus may further comprise a non-silent insertion, substitution, or deletion of a nucleotide sequence in at least one essential gene of the herpes simplex virus. In one embodiment, the herpes simplex virus lacks one expressible $\gamma_{134.5}$ gene, a non-essential gene. In a further embodiment, the recombinant herpes simplex lacks both expressible $\gamma_{134.5}$ genes.

In a preferred embodiment the cardiovascular condition is hypertension in a vascular tissue. Preferably, the cardiovascular condition is selected from the group consisting of chronic heart failure, hypertensive cardiovascular disease, ischemic heart disease, arrhythmia, congenital heart disease, valvular heart disease or stenotic defect, cardiomyopathy, aneurysm, chronic venous insufficiency, peripheral arterial disease, or restenosis.

In an additional embodiment, the heterologous nucleic acid sequence is expressed in vascular tissue for a duration selected from the group consisting of more than 7 days, more than 14 days, more than 21 days, more than 28 days, more than 35 days, or more than 70 days.

A further embodiment calls for the heterologous nucleic acid sequence to encode a screenable or selectable marker. Preferably, the heterologous nucleic acid sequence is an antithrombotic nucleic acid, an angiogenesis regulating nucleic acid, an immunomodulator, an inducer of cellular proliferation, an inhibitor of cellular proliferation or a regulator of programmed cell death.

In a related embodiment, the method of the present invention further comprises administering at least one therapeutic or pharmacological agent to the vascular cell. In a preferred embodiment, the pharmacological agent is selected from the group consisting of an antihyperlipoproteinemic agent, an

antiarteriosclerotic agent, an antithrombotic/fibrinolytic agent, a blood coagulant, an antiarrhythmic agent, an antihypertensive agent, a vasopressor, a treatment agent for congestive heart failure, an antianginal agent, and an anti-infection agent. Preferably, the recombinant herpes simplex virus is HSV-1. In a related embodiment, the recombinant herpes simplex virus further comprises at least one gene essential for the treatment of a herpes simplex virus infection by an anti-viral agent.

In yet another aspect of the present invention a method is provided for inducing normal physiology in a functionally abnormal vascular cell comprising administering to the cell a recombinant replicating herpes simplex viral vector comprising a heterologous nucleic acid sequence, and wherein the herpes simplex virus is debilitated for growth in the central nervous system.

The recombinant herpes simplex virus is debilitated for growth via non-silent insertion, substitution, or deletion of a nucleotide sequence in at least one non-essential gene of the herpes simplex virus. The recombinant herpes simplex virus may further comprise a non-silent insertion, substitution, or deletion of a nucleotide sequence in at least one essential gene of the herpes simplex virus.

Other features and advantages of the present invention will be better understood by reference to the detailed description and examples that follow.

DETAILED DESCRIPTION OF THE INVENTION

Somatic gene transfer has been proposed as a method to treat a variety of ischemic, thrombotic, and proliferative vascular disorders. However, progress in the field of vascular gene transfer has been hampered by a lack of vectors that can effectively penetrate the endothelium and stably and durably transform, transfect, or infect deeper cell layers.

Recently, the neurovirulence of herpes simplex virus type one (HSV-1) has been genetically attenuated, resulting in significantly reduced toxicity, while still maintaining its ability to penetrate and establish long-term infections. Deletion of both copies of the $\gamma 34.5$ gene potentially decreases virulence, but still allows for replication in dividing cells and, possibly, stable episomal infection. The product of

the γ 134.5 gene blocks phosphorylation of the α subunit of the translation initiation factor eIF-2 involved in viral protein synthesis.

To overcome the deficiencies in the art, a modified recombinant form of HSV-1 wherein both copies of the γ 134.5 gene were deleted (Chou et al., 1990; Chou and Roizman, 1994) was evaluated for its reliability and efficiency in transfecting vascular tissue *in vitro* and *in vivo*. The HSV-1 vector and methods of use are shown herein to be highly effective for gene transfer into vascular cells, with transfection efficiency and duration far exceeding that of traditional viral vectors. The disclosures herein demonstrate that a γ 134.5-deleted HSV-1 vector was capable of maintaining transgene expression in 50% of arterialized venous neointima for up to four weeks in animals with an intact immune system. These results were both surprising and unexpected, as the duration and efficiency demonstrated herein far exceeds the reported results with other vectors (Eslami et al., 2000; Schwartz et al., 1999; Scheinman et al., 1999; George et al., 2000; Dollery et al., 1999; Waugh et al., 2000; Hanna et al., 2000; Schneider et al., 1999). Further surprising and unexpected properties of the vector and methods of the present invention were demonstrated by the low dose of HSV-1 required to achieve these effects. Only 10^8 plaque forming units (pfu) per ml were used to achieve this duration and efficiency, which is a dose several orders of magnitude less than traditional strategies using adenovirus (Chang et al., 1995; Smith et al., 1997; Schwartz et al., 1999; Chang et al., 1995; Scheinman et al., 1999; Ueno et al., 1997; Cheng et al., 1998; George et al., 2000; Dollery et al., 1999; Bishop et al., 1999; Zoldhelyi et al., 2000; Waugh et al., 2000; Varenne et al., 1998; Cable et al., 1999; Fulton et al., 1996; Hanna et al., 1997; Neschis et al., 1998; Hanna et al., 2000; Schneider et al., 1999). Additionally, there was no apparent toxicity or immunostimulation at this low dose. However, the apparent lack of or reduced toxicity also was surprising and unexpected, given the propensity of HSV infection to cause arterial thrombosis in experimental systems (Key et al., 1990), as well as the indication that HSV may actually be involved in atherogenesis [Alber et al., *Circ.* 102:779-785, 2000].

In certain embodiments, HSV-1 vectors and methods of use described herein possess high efficiency (*e.g.*, greater than 40% transfection of cells) at low concentrations (*e.g.*, less than about 10^9 pfu per ml of administered vector). In

particular, the HSV-1 vectors described herein allow for reduced viral loads (*e.g.*, a total viral load of less than about 10^{10} pfu per ml).

In some embodiments, the HSV-1 vectors and methods of use described herein demonstrate long duration (*e.g.*, greater than about 28 days) of transgene
5 expression in a host cell, tissue, or organism. In particular aspects, the transgene duration is prolonged in despite the fact that the host organism is immunocompetent.

In particular embodiments, the HSV-1 vectors and methods of use described herein demonstrate a high level of cell penetrance (*e.g.*, greater than about 40%).

In certain embodiments, the HSV-1 vectors and methods of use described
10 herein demonstrate low or acceptable cytotoxicity (*e.g.*, less than about 10% cytotoxicity at a multiplicity of infection (MOI) ratio expressed as the total viral particles per cell of 2).

A. NUCLEIC ACIDS

Certain embodiments of the present invention concern various nucleic acids,
15 including vectors, promoters, therapeutic nucleic acids, and other nucleic acid elements that are involved in transformation and expression in cells. In certain aspects, a nucleic acid comprises a wild-type or a mutant nucleic acid. In particular aspects, a nucleic acid encodes a polypeptide or comprises a transcribed nucleic acid.

The term "nucleic acid" is well known in the art. A "nucleic acid" as used
20 herein will generally refer to a molecule (*i.e.*, a strand) of DNA, RNA or a derivative or analog thereof, comprising a nucleotide. A nucleotide includes, for example, a naturally occurring nucleotide containing a purine or pyrimidine base found in DNA (*e.g.*, an adenine "A," a guanine "G," a thymine "T" or a cytosine
25 "C") or RNA (*e.g.*, an A, a G, an uracil "U" or a C). The term "nucleic acid" encompass the terms "oligonucleotide" and "polynucleotide," each as a subgenus of the term "nucleic acid." The term "oligonucleotide" refers to a molecule of between about 3 and about 100 nucleotides in length. The term "polynucleotide" refers to at least one molecule of greater than about 100 nucleotides in length.

30 These definitions generally refer to a single-stranded molecule, but in specific embodiments will also encompass an additional strand that is partially, substantially, or fully complementary to the single-stranded molecule. Thus, a

nucleic acid may encompass a double-stranded molecule or a triple-stranded molecule that comprises one or more complementary strand(s) or "complement(s)" of a particular sequence comprising a molecule. As used herein, a single-stranded nucleic acid may be denoted by the prefix "ss", a double-stranded nucleic acid by the prefix "ds", and a triple-stranded nucleic acid by the prefix "ts."

In particular aspects, a nucleic acid encodes an RNA and/or a protein, polypeptide, and/or peptide. In certain embodiments, the present invention concerns novel compositions comprising at least one proteinaceous molecule. As used herein, a "proteinaceous molecule," "proteinaceous composition," "proteinaceous compound," "proteinaceous chain," or "proteinaceous material" generally refers, but is not limited to, a protein of greater than about 200 amino acids or the full length endogenous sequence translated from a gene; a polypeptide of greater than about 100 amino acids; and/or a peptide of from about 3 to about 100 amino acids. All the "proteinaceous" terms described above are used interchangeably herein.

A.1. PREPARATION OF NUCLEIC ACIDS

A nucleic acid may be made by any technique known to one of ordinary skill in the art, such as, for example, chemical synthesis, enzymatic production or biological production. Non-limiting examples of a synthetic nucleic acid (*e.g.*, a synthetic oligonucleotide), include a nucleic acid made by *in vitro* chemical synthesis using phosphotriester, phosphite, or phosphoramidite chemistry and solid-phase techniques such as described in EP 266,032, incorporated herein by reference, or via deoxynucleoside H-phosphonate intermediates as described by Froehler et al., 1986 and U.S. Patent Serial No. 5,705,629, each incorporated herein by reference. In the methods of the present invention, one or more oligonucleotides may be used. Various different mechanisms of oligonucleotide synthesis have been disclosed in, for example, U.S. Patents. 4,659,774, 4,816,571, 5,141,813, 5,264,566, 4,959,463, 5,428,148, 5,554,744, 5,574,146, 5,602,244, each of which is incorporated herein by reference.

A non-limiting example of an enzymatically produced nucleic acid includes one produced by enzymes in amplification reactions such as PCRTM (see, for example, U.S. Patent 4,683,202 and U.S. Patent 4,682,195, each incorporated

herein by reference), or the synthesis of an oligonucleotide described in U.S. Patent No. 5,645,897, incorporated herein by reference. A non-limiting example of a biologically produced nucleic acid includes a recombinant nucleic acid produced (*i.e.*, replicated) in a living cell, such as a recombinant DNA vector replicated in bacteria (see for example, Sambrook *et al.* 1989).

A.2. PURIFICATION OF NUCLEIC ACIDS

A nucleic acid may be purified on polyacrylamide gels, cesium chloride centrifugation gradients, or by any other means known to one of ordinary skill in the art (see for example, Sambrook *et al.*, 1989. In preferred aspects, a nucleic acid is a pharmacologically acceptable nucleic acid. Pharmacologically acceptable compositions are known to those of skill in the art; exemplars are described herein.

In some embodiments, the present invention concerns a nucleic acid that is an isolated nucleic acid. As used herein, the term "isolated nucleic acid" refers to a nucleic acid molecule (*e.g.*, an RNA or DNA molecule) that has been isolated free of, or is otherwise free of, the bulk of the total genomic and transcribed nucleic acids of one or more cells. In certain embodiments, "isolated nucleic acid" refers to a nucleic acid that has been isolated free of, or is otherwise free of the bulk of cellular components or *in vitro* reaction components such as, for example, macromolecules such as lipids or proteins, small biological molecules, and the like.

A.3. NUCLEIC ACID SEGMENTS

In certain embodiments, the nucleic acid is a nucleic acid segment. As used herein, the term "nucleic acid segment," is a smaller fragment of a nucleic acid, such as, for non-limiting example, those that encode only part of a peptide or polypeptide sequence. Thus, a "nucleic acid segment" may comprise any part of a gene sequence, of from about 2 nucleotides, and preferably 50 nucleotides, to the full length of a peptide or polypeptide encoding region.

Various nucleic acid segments may be designed based on a particular nucleic acid sequence, and may be of any length. By assigning numeric values to a sequence, for example, the first residue is 1, the second residue is 2, etc., an algorithm defining all nucleic acid segments can be created:

5 n to $n + y$

where n is an integer from 1 to the last number of the sequence and y is the length of the nucleic acid segment minus one, where $n + y$ does not exceed the last number of the sequence. Thus, for a 10-mer, the nucleic acid segments correspond to bases 1 to 10, 2 to 11, 3 to 12 ... and so on. For a 15-mer, the nucleic acid segments correspond to bases 1 to 15, 2 to 16, 3 to 17, and so on. For a 20-mer, the nucleic segments correspond to bases 1 to 20, 2 to 21, 3 to 22, and so on. In certain embodiments, the nucleic acid segment may be a probe or primer. As used herein, a "probe" generally refers to a nucleic acid used in a detection method or composition. As used herein, a "primer" generally refers to a nucleic acid used in an extension or amplification method or composition.

15 A.4. NUCLEIC ACID COMPLEMENTS

The present invention also encompasses a nucleic acid that is complementary to a nucleic acid. A nucleic acid "complement(s)" or is "complementary" to another nucleic acid when it is capable of base-pairing with another nucleic acid according to the standard Watson-Crick, Hoogsteen, or reverse Hoogsteen complementarity rules. As used herein "another nucleic acid" may refer to a separate molecule or a spatially separated sequence of the same molecule. In preferred embodiments, a complement is an antisense nucleic acid used to reduce expression (*e.g.*, translation) of a RNA transcript *in vivo*.

25 As used herein, the term "complementary" or "complement(s)" also refers to a nucleic acid comprising a sequence of consecutive nucleotides or semi-consecutive nucleotides (*e.g.*, one or more of the consecutive nucleotides is not present in the molecule) capable of hybridizing to another nucleic acid strand or duplex even if less than all the nucleotides do not base pair with a counterpart nucleotide. However, in some antisense embodiments, completely complementary nucleic acids are preferred.

B. VECTORS

The term "vector" is used to refer to a carrier nucleic acid molecule into which a nucleic acid sequence can be inserted for introduction into a cell where it can be replicated. As used herein, "vector" refers to both the carrier molecule and a recombinant molecule containing an insert (typically heterologous) nucleic acid and the carrier molecule. The usages will be apparent to one of skill from the context of a recitation. A nucleic acid segment can be "exogenous," which means that it is not a sequence originally from the genome of the cell into which the segment is being introduced. As used herein, a "heterologous" nucleic acid is defined herein to mean a nucleic acid segment that is foreign with respect to another nucleic acid sequence. For example, a nucleic acid segment may be heterologous to the genome of a cell or to another juxtaposed nucleic acid segment that together comprise a larger nucleic acid with the first nucleic acid segment.

The term "expression vector" refers to any type of genetic construct comprising a nucleic acid coding for a RNA capable of being transcribed. In some cases, RNA molecules are then translated into a protein, polypeptide, or peptide. In other cases, these sequences are not translated, for example, in the production of antisense molecules or ribozymes. Expression vectors can contain a variety of "control sequences," which refer to nucleic acid sequences necessary for the transcription and possibly translation of an operably linked coding sequence in a particular host cell. In addition to control sequences that govern transcription and translation, vectors and expression vectors may contain nucleic acid sequences that serve other functions as well and are described below.

As referred to herein and at a minimum, expression of a heterologous nucleic acid sequence is defined as expression of a cognate RNA molecule (typically an mRNA or rRNA). More generally, expression of a heterologous nucleic acid sequence refers to the cellular processes of transcription and translation that produce a polypeptide gene product. Although the polypeptide gene product would typically include at least one function characteristic of that polypeptide, as referred to herein, expression of a heterologous nucleic acid sequence is further contemplated to include the product of a non-functional polypeptide.

B.1. HSV VIRAL VECTORS

Herpes simplex virus (HSV) type I and type II contain a double-stranded, linear DNA genome of approximately 150 kb, containing at least 84 expressible open reading frames [see Table 1, which sets forth the function of herpes simplex genes (Roizman, *Proc. Natl. Acad. Sci.* 93:11307-11312, October 1996)]. Wild-type HSVs are able to infect cells lytically and to establish latency in certain cell types (*e.g.*, neurons). Similar to adenovirus, HSV also can infect a variety of cell types including muscle (Yeung et al., 1999), ear (Derby et al., 1999), eye (Kaufman et al., 1999), tumors (Yoon et al., 1999; Howard et al., 1999), lung (Kohut et al., 1998), neuronal (Garrido et al., 1999; Lachmann and Efsthathiou, 1999), liver (Miyake et al., 1999; Kooby et al., 1999) and pancreatic islets (Rabinovitch et al., 1999).

HSV viral genes are transcribed by cellular RNA polymerase II and are temporally regulated, resulting in the transcription and subsequent synthesis of gene products in roughly three discernable phases or kinetic classes. These phases of genes are referred to as the Immediate Early (IE) or α genes, Early (E) or β genes and Late (L) or γ genes. Immediately following the arrival of the genome of a virus in the nucleus of a newly infected cell, the IE genes are transcribed. The efficient expression of these genes does not require prior viral protein synthesis. The products of IE genes are required to activate transcription and regulate the remainder of the viral genome.

HSV can infect a wide variety of both dividing and non-dividing host cells, and thus has been used in the production of herpes virus vectors for delivery of heterologous nucleic acids into host cells. In certain embodiments, a herpes simplex virus vector may be a HSV-1 or HSV-2 vector. Particularly preferred vectors are viral vectors that lack one or more copies of a $\gamma_{134.5}$ gene, or ribonucleotide reductase gene (*i.e.*, the γ_{139} gene encoding ICP6 or the γ_{140} gene). Vectors containing a single copy of the $\gamma_{134.5}$ gene lack the inverted repeat associated with virulence. Viral vectors can be mutated by the methods disclosed herein or as would be known to those of skill in the art to produce a viral vector that lacks at least one $\gamma_{134.5}$ gene and/or ribonucleotide reductase gene for use in the methods disclosed herein. For example, methods of producing herpes virus vectors that lack

one or more γ 134.5 genes or ribosomal reductase genes are disclosed in U.S. Patent Nos. 6,120,773, 6,106,826, 5,795,713, 5,585,096 and 5,328,688, each incorporated herein by reference.

Embodiments would include use (in the methods of the present invention) of herpes simplex viral vectors that are debilitated for growth within the central nervous system. As used herein, a debilitated HSV is defined in the following manner. A wild-type strain (non-recombinant) requires 10 pfu to 1000 pfu to kill a mouse by intracerebral inoculation. A debilitated virus requires at least 100,000 pfu to kill a mouse host. Further, a virulent virus travels from cell to cell in a rapid fashion, while a debilitated virus is highly restricted and does not travel from cell to cell very efficiently (hence more virus particles are required to kill a host such as a mouse), unless the animal host cannot immunologically respond to the infection.

HSV may be debilitated for growth in the CNS and useful in the present methods by a non-silent nucleotide insertion, substitution, or deletion in at least one non-essential gene of a herpes simplex virus. Examples of non-essential genes and their functions are set forth in Table 3. Such a debilitated HSV may alternatively comprise a non-silent nucleotide insertion, substitution, or deletion in at least one essential gene of a herpes simplex virus. Examples of essential genes and their functions are shown in Table 2

As used herein an essential gene is defined as a gene that is required to produce infectious progeny in a cell. For example, the HSV gene, α 4, is essential, *i.e.*, in its absence the virus will not produce infectious progeny. However, one of skill in the art would recognize that a cell line producing the α 4 gene product will allow a deletion mutant (α 4⁻) to replicate. As used herein a non-essential gene is defined as a gene that, when absent or dysfunctional, does not prevent the virus from replicating and producing progeny in a cell line that does not supply the missing gene product. For example, the HSV gene, α 0, is considered non-essential, *i.e.*, an α 0 deletion mutant (α 0⁻) produces progeny in many different cell lines (in some cell lines the titer is 50-100 fold lower, while in other cell lines nearly normal yields are produced.

To ensure stable modification of genes such as γ 134.5, which are located within the inverted repeat of HSV, the inverted repeat itself is modified sufficiently

to prevent reconstitution at the unmodified gene. Without wishing to be bound by theory, it is believed that gene reconstitution, or gene conversion occurs as a result of inverted repeat-mediated pairing of duplicated genes found partially or wholly within the repeat region, with one copy of a gene (*e.g.*, the unmodified copy) serving as a template for conversion of the other copy of the gene (*e.g.*, the modified copy). Accordingly, one of skill, using routine procedures, would modify an inverted repeat in addition to a gene contained therein, sufficiently to prevent the effective loss of the modified gene. For example, one of skill would modify the sequence of one inverted repeat sufficiently to prevent effective *in vivo* pairing of the repeats and genes they contain, which is expected to prevent the loss of a modified gene found in the repeat, believed to be due to the prevention of gene conversion.

Table 1

GENE	Product	Dispensable in cell culture	Regulation	Function of gene product
<i>γ34.5</i>	ICP34.5	Y	γ_1	Null mutants are attenuated and fail to block phosphorylation of eIF-2 α by activated protein kinase; RNA-dependent kinase; carboxyl terminus homologous to the corresponding domain of the GADD34 proteins.
<i>ORF-P</i>	ORF-P	Y	pre α	ORF is antisense to the γ 34.5 gene and repressed by binding of ICP4 to cap site. Proteins interact with p32, a component of SF2/ASF splicing factor.
<i>ORF-O</i>	ORF-O	Y	pre α	Overlaps with ORF-P, a protein made by frameshift from ORF-P.
<i>α0</i>	ICP0	Y	α	Promiscuous transactivator, requires ICP4 for optimal activity; nucleotidylated, phosphorylated by UL13, nuclear (early) and cytoplasmic (late) phases. Null mutants debilitated at low multiplicities of infection.
<i>UL1</i>	gL	N	γ	Complex with gH required for transport of both proteins to plasma membrane and for viral entry mediated by gH.
<i>UL2</i>		Y	β	Uracil DNA glycosylase.
<i>UL3</i>		Y	γ_2	Nuclear phosphoprotein of unknown function. Reported to localize to perinuclear region early and to the nucleus late in infection.
<i>UL4</i>		Y	Unknown	Unknown.
<i>UL5</i>		N	β	Forms complex with UL8 and UL52 proteins.
<i>UL6</i>		N	Unknown	Virion protein; required for DNA cleavage and packaging.
<i>UL7</i>		N	Unknown	Unknown.
<i>UL8</i>		N	β	Forms complex with UL5 and UL52 (helicase/primase complex). Stabilizes interaction between primers and DNA template.
<i>UL9</i>		N	$\gamma(?)$	Binds to origins of DNA synthesis in sequence-specific (origin) fashion; carries out helicase and ATPase activities.
<i>UL10</i>	gM	Y	γ	Glycoprotein present in virions and plasma membranes.

Table 1 Continued

UL10.5		?	Unknown	Unknown.
UL11		Y	γ (?)	Myristoylated protein; necessary for efficient capsid envelopment and exocytosis.
UL12		Y	β	Exonuclease (DNase) involved in viral nucleic acid metabolism; reported to localize in nucleoli and in virally induced nuclear dense bodies and to bind to a sequence along with other unidentified proteins. Complex may be involved in cleavage/packaging of viral DNA.
UL12.5		Y	Unknown	Nuclease-associated with capsids.
UL13		Y	γ	Virion (nuclear) protein kinase; substrates include ICP0, ICP22, <i>whs</i> , UL3, UL49, etc.
UL14		N	Unknown	Unknown.
UL15		N	γ	<i>ts</i> mutant DNA +. Two exons; protein required for cleavage/packaging of DNA.
UL16		Y	Unknown	Virion protein; gene located within intron of UL15.
UL17		N	γ	Located within intron of UL15.
UL18	VP23	N	γ	Protein required for capsid formation and cleavage/packaging of DNA.
UL19	VP5, ICP5	N	γ_1	Major capsid protein.
UL20		Y	γ	Membrane protein, associates with nuclear membranes, Golgi stacks, etc. Essential for viral exocytosis.
UL20.5			γ_2	Unknown.
UL21		Y	Unknown	Nucleotidylated phosphoprotein; unknown function.
UL22	gH	N	γ_2	Forms complex with gL (see above). Required for entry, egress, and cell-cell spread.
UL23	ICP36	Y	β	Thymidine (nucleoside) kinase.
UL24		Y	γ	Syn' locus; membrane-associated protein?
UL25		N	γ	Virion protein reported to be required for packaging of cleaved viral DNA.
UL26		N	γ	Serine protease; substrates are UL26 protein and UL26.5 (ICP35). (C portion of UL26), VP24 (N terminus of protease) are products of the self-cleavage of UL26.
UL26.5	ICP35	N	γ	Substrate of UL26 protease unique to B capsids and forms inner core or scaffolding; the precursor, ICP35b,c is cleaved to e, f. On packaging of DNA it is removed from capsid shell.

Table 1 Continued

U _L 27	gB, VP7	N	γ^1	Glycoprotein required for viral entry; forms a dimer and induces neutralizing antibody. A syn locus maps to the carboxyl terminus.
U _L 27.5		?	Unknown	Unknown, antisense to gB.
U _L 28	ICP18.5	N	γ	M _r 87-95 K protein required for DNA cleavage/packaging.
U _L 29	ICP8	N	β	Binds single-stranded DNA cooperatively, required for viral DNA replication: forms complex with DNA polymerase and U _L 42. <i>ts</i> mutants are DNA ⁻ and hence expression of early and late genes may be affected positively or negatively by ICP8. Because ICP8 denatures DNA, it affects renaturation of complementary strands of DNA and affects homologous pairing and strand transfer.
U _L 30		N	β	DNA polymerase; forms complex with ICP8 and C terminal 247 amino acids of U _L 42.
U _L 31		N	γ^2	Nucleotidylated phosphoprotein, cofractionates with nuclear matrix.
U _L 32		N	γ^2	Cytoplasmic/nuclear protein required for DNA cleavage/packaging.
U _L 33		N	Unknown	DNA packaging; necessary for assembly of capsids containing DNA.
U _L 34		N	Unknown	Abundant nonglycosylated, membrane-associated, virion protein phosphorylated by U _S 3.
U _L 35	VP26	N	γ^2	Basic phosphorylated capsid protein.
U _L 36	ICP1-2	N	γ^2	Tegument phosphoprotein. DNA is not released from capsids at nuclear pores in cells infected with <i>ts</i> mutant. Reported to form complex with a M _r 140 K protein that binds <i>a</i> sequence DNA.
U _L 37	ICP32	N	γ	Cytoplasmic phosphoprotein; in presence of ICP8 it is transported to nucleus and associates with DNA, but phosphorylation is not dependent on ICP8. Required for maturation of virions.
U _L 38	VP19C	N	γ^2	Capsid assembly protein, blinds DNA and may be involved in anchoring DNA in the capsid.
U _L 39	ICP6	Y	β	Large subunit of ribonucleotide reductase. Autophosphorylates via unique N terminus but does not trans-phosphorylate.
U _L 40		Y	β	Small subunit of ribonucleotide reductase.
U _L 41	VHS	Y	γ	Causes nonspecific degradation of mRNA after infection; shuts off host protein synthesis, enables sequential synthesis of viral proteins.
U _L 42		N	β	Double-stranded DNA-binding protein, binds to and increases processivity of DNA polymerase.

Table 1 Continued

UL43		Y	Unknown	Amino acid sequence predicts membrane-associated protein.
UL43.5		Y		Antisense to UL43; low abundance nuclear protein; accumulates in assemblons.
UL44	gC, vp7.5	Y	γ_2	Glycoprotein involved in cell attachment; required for attachment to the apical surface of polarized MDCK cells.
UL45		Y	γ_2	Encodes a Mr 18 K protein of unknown function.
UL46	VP11/12	Y	γ	Tegument phosphoprotein reported to modulate the activity of UL48 (aTIF).
UL47	VP13/14	Y	γ_2	Nucleotidylated tegument phosphoprotein modulates the activity of UL48 (aTIF).
UL48	VP16, aTIF	N	γ	Tegument protein, induces γ genes by interacting with <i>Ocr1</i> . The complex binds to specific sequences with the consensus GyATGnTAATGArATTCTTGGG-NC.
UL49	VP22	N	γ	Nucleotidylated, mono(ADP-ribosyl)ated tegument phosphoprotein.
UL49.5		N	γ_2	Sequence predicts a Mr 12,000 membrane-associated protein.
UL50		Y	β	dUTPase.
UL51		Y	γ	Unknown.
UL52		N	β	Component of the helicase/primase complex.
UL53	gK	Y	γ	Glycoprotein required for efficient viral exocytosis; contains <i>syn</i> locus.
$\alpha 27$	ICP27	N	α	Nucleotidylated multifunctional regulatory protein; causes redistribution of snRNPs, inhibits RNA splicing. It is required for late gene expression, and negatively regulates early genes.
UL55		Y	Unknown	Unknown.
UL56		Y	Unknown	Nuclear, virion-associated protein of unknown function.
$\alpha 4$	ICP4	N	α	Nucleotidylated, poly(ADP-ribosyl)ated phosphoprotein; regulates positively most β and γ genes and negatively itself, ORF-P and the $\alpha 0$ gene; blocks apoptosis. Binds to DNA in sequence specific fashion.
$\alpha 22$	ICP22	Y	α	Nucleotidylated regulatory protein, phosphorylated by UL13 and Us3 protein kinases, required for optimal expression of ICP0 and of a subset of γ proteins.
Us1.5	Us1.5	Y	α	Regulatory protein; extent to which it shares function with ICP22 not known.

Table 1 Continued

Us2		Y	Unknown	Unknown.
Us3		Y	β	Protein kinase; major substrate is UL34 protein.
Us4	gG	Y	γ	Glycoprotein involved in entry, egress, and spread from cell to cell.
Us5	gL(?)	Y	Unknown	Sequence predicts glycoprotein.
Us6	gD VP17/18	N	γ_1	Glycoprotein required for post-attachment entry of virus into cells.
Us7	gL	Y	γ	gL and gE glycoproteins form complex for transport to plasma membrane and also to constitute a high-affinity Fc receptor. GL is required for basolateral spread of virus in polarized cells.
Us8	gE	Y	γ_2	FC receptor; involved in basolateral spread of virus in polarized cells.
Us8.5		Y	β or γ_1	Unknown.
Us9		Y	Unknown	Tegument protein phosphorylated by UL13.
Us10		Y	Unknown	Tegument protein.
Us11		Y	γ_2	Tegument protein binds to UL34 mRNA in sequence- and conformation-specific fashion; binds to the 60S ribosomal subunit and localizes in the nucleolus.
cd47	ICP47	Y	α	Binds to TAP1/TAP2 and to block antigen presentation to CD8+ cells.
OrisTU	OrisRNA	Y	γ_2	RNA transcribed across S origins of DNA synthesis. Function is not known.
LATU	LATs	Y	pre α ?	Transcripts, found in latently infected neurons. Function is not known.

Table 2

EXEMPLARY ESSENTIAL HSV GENES		
Group	Number of Genes	Function (Genes)
1	4	Virus attachment, penetration, envelopment, and exocytosis, (gB, gD, gH, gL)
2	2	Regulation of gene expression: $\alpha 4$: positive and negative control of gene expression $\alpha 27$: post transcriptional processing, transport of RNA.
3	7	Replication of viral DNA (UL5, 8, 9, 29, 30, 42, 52)
4	8	Capsid proteins including the protease (UL6, 17, 18, 19, 26, 26.5, 35, 38)
5	6	Packaging of DNA (UL15, 25, 28, 31, 32, 33)
6	10	Virion assembly and other functions (UL7, 14, 27.5, 34, 36-38, 48, 49, 49.5)

Table 3

EXEMPLARY NON-ESSENTIAL HSV GENES		
Group	Number of Genes	Function (Genes)
1	12	Entry into polarized cells, sorting, exocytosis: (gC, gE, gG, gI, gJ, gK, gM, UL11, 20, 24, 43, 45)
2	4	Block host response to infection: γ 34.5 blocks phosphorylation of eIF-2 α α 47: blocks presentation of antigenic peptides to the immune system. Us9: binds ubiquitin and proteasomes. Us3: protein kinase, blocks apoptosis.
3	7	Enable viral replication in nondividing cells: thymidine kinase, dUTPase, DNase ribonucleotide reductase, uracil glycosylase
4	1	Degrade cellular mRNA and facilitated expression of viral genes. UL41
5	18	Functions are not known: Us2, 8.5, 10; UL3, 4, 8.5, 9.5, 13, 16, 20.5, 21, 43.5, 46, 47, 51, 55, 56; ICP 22
6	2	Genes suppressed during productive infection: ORF P: inhibits synthesis of proteins from spliced RNA. ORF O: blocks the ICP4 transactivator from binding DNA.

As used herein, a viral vector "lacks" a gene when it is mutated so that one or more copies of the mutated gene possess reduced or absent expression or activity sufficient to debilitate it for growth in the central nervous system (CNS) or neurons as defined above. Such mutations may comprise, for example, frame-shift mutations, insertions, deletions, truncations, and the like in coding or regulatory sequences for one or more genes. The product of the $\gamma_134.5$ gene contributes to neurovirulence (Chou et al., 1994). A viral vector that lacks one or more $\gamma_134.5$ genes has attenuated neurovirulence, but does not block protein synthesis and will cause the death of normal host cells (of course, deletion of the $\gamma_134.5$ gene typically deletes part or all of the overlapping and oppositely oriented orfp. HSV viral vectors that lack one or more ribonucleotide reductase genes also have attenuated neurovirulence (*i.e.*, loss of either subunit of ribonucleotide reductase results in attenuation) and are susceptible to antiviral drugs such as acyclovir and ganciclovir, allowing for antiviral therapy in instances wherein a host exhibits an undesired side effect of vector administration [*e.g.*, viral encephalitis, U.S. Patent 6,106,826]. It is contemplated that one or more other genes in a herpes viral vector may be mutated to improve the efficacy or reduce the side effects of a herpes viral vector. Such genes are known in the art, and include the thymidine kinase (tk) gene, or other genes required for HSV replication, infection or latency.

For use in therapeutic gene delivery, HSV may be replication-defective. Protocols for generating replication-defective HSV helper virus-free cell lines have been described (U. S. Patent 5,879,934; U. S. Patent 5,851,826, each specifically incorporated herein by reference in its entirety). One IE protein, Infected Cell Polypeptide 4 (ICP4), encoded by the α_4 gene, is absolutely required for both virus infectivity and the transition from IE to later transcription. Thus, due to its complex, multifunctional nature and central role in the regulation of HSV gene expression, ICP4 has typically been the target of HSV genetic studies.

Phenotypic studies of HSV viruses deleted for ICP4 indicate that such viruses are potentially useful for gene transfer purposes (Krisky et al., 1998a). One property of viruses deleted for ICP4 that makes them desirable for gene transfer is that they only express the five other IE genes: U_s 1.5, ICP0, ICP27, ICP22 and ICP47 (DeLuca et al., 1985) without the expression of viral genes encoding proteins

that direct viral DNA synthesis, as well as the structural proteins of the virus. This property is desirable for minimizing possible deleterious effects on host cell metabolism or an immune response following gene transfer. Further deletion of IE genes ICP22 and ICP27, in addition to ICP4, substantially improve reduction of HSV cytotoxicity and prevent early and late viral gene expression (Krisky et al., 1998b).

The therapeutic potential of HSV in gene transfer has been demonstrated in various *in vitro* model systems and *in vivo* for diseases such as Parkinson's (Yamada et al., 1999), retinoblastoma (Hayashi et al., 1999), intracerebral and intradermal tumors (Moriuchi et al., 1998), B cell malignancies (Suzuki et al., 1998), ovarian cancer (Wang et al., 1998) and Duchenne Muscular Dystrophy (Huard et al., 1997).

Thus, in some embodiments, a viral vector comprises a nucleic acid sequence that encodes a therapeutic nucleic acid. In preferred aspects, a therapeutic nucleic acid encodes antisense nucleic acid or proteinaceous composition effective to treat a vascular, cardiac, and/or cardiovascular disease or disorder. One of skill in the art would be well equipped to construct a vector through standard recombinant techniques (see, for example, Maniatis et al., 1988 and Ausubel et al., 1994. Non-limiting examples of virus vectors that may be used to deliver a nucleic acid of the present invention are described below.

B.2. THERAPEUTIC NUCLEIC ACIDS

In certain embodiments, a vector of the present invention comprises heterologous nucleic acid. In preferred aspects, a heterologous nucleic acid comprises a therapeutic nucleic acid encoding a proteinaceous composition or an antisense product. Exemplary heterologous nucleic acids encode a nucleotide (*e.g.*, antisense), protein, polypeptide or peptide product having an antiproliferative, vasodilatory or angiogenic activity.

The nucleotide and protein, polypeptide and peptide sequences for various genes have been previously disclosed, and may be found at computerized databases known to those of ordinary skill in the art. One such database is the National Center for Biotechnology Information's Genbank and GenPept databases. The coding regions for these known genes may be amplified, cloned into and/or expressed in the vectors of the present invention using the techniques disclosed

herein or by any technique that would be known to those of ordinary skill in the art (see, for example, Sambrook et al., 1989).

In certain embodiments, a vector may comprise more than one such therapeutic nucleic acid. The therapeutic nucleic acids could be from the same functional group (*e.g.*, both antiproliferative, vasodilatory or angiogenic, etc.) or from different functional groups (*e.g.*, an antiproliferative and an angiogenic). By presenting particular combinations of therapeutic nucleic acids to a target cell, it may be possible to augment the overall effect of either or both nucleic acids or their encoded products on the physiology of the target cell.

For example, deposits of extracellular matrix proteins and vascular cell proliferation are associated with atherosclerosis. Genes such as acidic fibroblast growth factor (aFGF) that are overexpressed in, *e.g.*, porcine arteries result in smooth muscle cell proliferation and resultant arterial wall thickening, as well as new blood vessel formation in the arterial wall due to endothelial cell migration (Nabel et al, 1993). TGF- β 1 and platelet-derived growth factor B (PDGFB) overexpression have also led to vessel wall thickening. In certain aspects, it is contemplated that expression, in vascular or cardiovascular cells, of nucleic acids that bind RNAs encoding proteins that contribute to atherosclerosis (*e.g.*, aFGF, TGF- β 1, PDGFB) will reduce the progress of atherosclerosis.

In another example, it has been demonstrated that administration of a vector that expresses a nucleic acid whose encoded product inhibits cell growth can inhibit restenosis in a patient. For example, an adenoviral vector that expressed thymidine kinase, which makes cells susceptible to killing by ganciclovir, coupled with administration of ganciclovir, prevented smooth muscle proliferation and restenosis in an animal model system after balloon angioplasty (Ohno et al., 1994). Thus, in particular facets, it is contemplated that administration of vectors that express such therapeutic nucleic acids for blood vessel thickening may be combined with administration of a pharmacological agent (*e.g.*, an antihyperlipoproteinemic, an antiarteriosclerotic, an antithrombotic/fibrinolytic agent, an antihypertensive agent) and/or a surgical procedure (*e.g.*, arterial bypass surgery, balloon angioplasty) used in the treatment of patients suffering from atherosclerosis or restenosis.

In preferred embodiments, the therapeutic nucleic acid will be effective in treating a vascular or cardiovascular disease or condition. A cardiovascular disease or condition which may benefit from administration of a vector of the present invention includes, but is not limited to, chronic heart failure, hypertensive cardiovascular disease, ischemic heart disease, arrhythmia, congenital heart disease, valvular heart disease or stenotic defect, cardiomyopathy, aneurysm, chronic venous insufficiency, peripheral arterial disease or a combination thereof.

In certain embodiments, a vector of the present invention may encode at least one antiproliferative, vasodilatory or angiogenic proteinaceous composition. For example, in certain aspects, a vector of the present invention may comprise an antithrombotic nucleic acid, a nucleic acid that regulates angiogenesis (*e.g.*, VEGF, aFGF, bFGF), an immunomodulatory nucleic acid (*e.g.*, angiostatin, endostatin, TNF α , IL-1, IL-10), a cell proliferation stimulatory nucleic acid (*e.g.*, platelet-derived growth factor, ras, c-myc, c-fos), a nucleic acid that encodes an inhibitor of cell proliferation (*e.g.*, p53, p21, retinoblastoma protein (Rb)) or antisense expressing constructs of such nucleic acids.

In further embodiments, a vector of the present invention may encode one or more enzymes, including, but not limited to, angiogenin, proliferating cell nuclear antigen (PCNA), senescent cell-derived inhibitor 1 (sdi1), a cell cycle division protein (cdc), a cyclin-dependent kinase (cdk), C-type natriuretic peptide (CNP); a homeobox gene including, but not limited to, gax, thymidine kinase (tk), cytosine deaminase, endothelial cell nitric oxide synthase (ec-NOS), inducible nitric oxide synthase (iNOS), hirudin; a cell-adhesion molecule including, but not limited to, VCAM, ICAM, PECAM or E-selectin; a tissue inhibitor of a metalloproteinase (TIMP); analogs to cAMP such as adrenomedulin, mitogen-activated protein kinases (MAPKs) such as extracellular response kinases (ERKs) and c-jun N-terminal kinase (JNK) and their inhibitors, and their antisense oligonucleotides or polypeptides. In other embodiments, a vector of the present invention may encode one or more polynucleotides. In other embodiments, a vector of the present invention may comprise an oncogene including, but not limited to, c-myb.

B.2.a. Antithrombotic Nucleic Acids

In certain embodiments, a therapeutic nucleic acid may produce an anti-thrombotic effect. Non-limiting examples of such nucleic acids include those that encode COX-1, TFPI, PGS, Dp, ras, myc, neu, raf, erb, fms, trk, ret, gsp, hst, 5
abl, E1A, hirudin, t-PA or p300 coding sequences.

B.2.b. Angiogenesis-Regulating Nucleic Acids

In other embodiments, a vector of the present invention may encode a nucleic acid that increases or inhibits angiogenesis. Non-limiting examples of nucleic acids that may regulate angiogenesis include vascular endothelial growth 10
factor (VEGF), acidic fibroblast growth factor (aFGF), basic fibroblast growth factor (bFGF), thrombospondin, BAI-1, GDAIF, TNF α , or their receptors.

B.2.c. Immunomodulators

It is contemplated that immunomodulators can be included in the vector to augment a cell's or a patient's (*e.g.*, an animal's) response. Immunomodulators can 15
be included as purified proteins, nucleic acids encoding immunomodulators, and/or cells that express immunomodulators in a vaccine composition. The following sections list non-limiting examples of immunomodulators that are of interest, and it is contemplated that various combinations of immunomodulators may be used in certain embodiments (*e.g.*, a cytokine and a chemokine).

B.2.c.i. Cytokines

Interleukins, cytokines, nucleic acids encoding interleukins or cytokines, and/or cells expressing such compounds are contemplated as heterologous nucleic acid gene products. Interleukins and cytokines, include, but are not limited to 25
interleukin 1 (IL-1), IL-2, IL-3, IL-4, IL-5, IL-6, IL-7, IL-8, IL-9, IL-10, IL-11, IL-12, IL-13, IL-14, IL-15, IL-18, β -interferon, α -interferon, γ -interferon, angiostatin, thrombospondin, endostatin, GM-CSF, G-CSF, M-CSF, METH-1, METH-2, tumor necrosis factor (TNF; Cachectin), TGF β , LT and combinations thereof.

B.2.c.ii. Chemokines

Chemokines, nucleic acids that encode for chemokines, and/or cells that 30
express such also may be used as heterologous nucleic acids. Chemokines generally act as chemoattractants to recruit immune effector cells to the site of chemokine

expression. It may be advantageous to express a particular chemokine coding sequence in combination with, for example, a cytokine coding sequence, to enhance the recruitment of other immune system components to a site of treatment, *e.g.*, in vascular tissue. Such chemokines include, for example, RANTES, MCAF, MCP-1, MIP1-alpha, IL-8, MIP1-Beta, IP-10 and combinations thereof. The skilled artisan will recognize that certain cytokines are also known to have chemoattractant effects and could also be classified as chemokines.

B.2.d. Inducers of Cellular Proliferation

In one embodiment of the present invention, it is contemplated that anti sense mRNA directed to a particular inducer of cellular proliferation is used to prevent expression of the inducer of cellular proliferation. The proteins that induce cellular proliferation further fall into various categories dependent on function. The commonality of all of these proteins is their ability to regulate cellular proliferation.

For example, a form of PDGF, the *sis* oncogene, is a secreted growth factor. The proteins FMS, ErbA, ErbB and *neu* are also growth factor receptors. Mutations to these receptors result in loss of regulatable function. For example, a point mutation affecting the transmembrane domain of the *Neu* receptor protein results in the *neu* oncogene. The *erbA* oncogene is derived from the intracellular receptor for thyroid hormone. The modified oncogenic ErbA receptor is believed to compete with the endogenous thyroid hormone receptor, causing uncontrolled growth.

The largest class of oncogenes includes the signal transducing proteins (*e.g.*, Src, Abl and Ras). The protein Src is a cytoplasmic protein tyrosine kinase, and its transformation from proto-oncogene to oncogene, in some cases, results via mutations at tyrosine residue 527. In contrast, transformation of the GTPase protein *ras* from proto-oncogene to oncogene, in one example, results from a valine to glycine mutation at amino acid 12 in the sequence, reducing *ras* GTPase activity.

Other proteins such as Jun, Fos and Myc are proteins that directly exert their effects on nuclear functions as transcription factors.

B.2.e. Inhibitors of Cellular Proliferation

In certain embodiment, the restoration of the activity of an inhibitor of cellular proliferation through a genetic construct is contemplated. Tumor

suppressor oncogenes function to inhibit excessive cellular proliferation. The inactivation of these genes destroys their inhibitory activity, resulting in unregulated proliferation. The tumor suppressors p53, p16, p21 and C-CAM are examples of this class of proteins.

5 High levels of mutant p53 have been found in many cells transformed by chemical carcinogenesis, ultraviolet radiation, and several viruses. The p53 gene is a frequent target of mutational inactivation in a wide variety of human tumors and is already documented to be the most frequently mutated gene in common human cancers. It is mutated in over 50% of human NSCLC (Hollstein et al., 1991) and in
10 a wide spectrum of other tumors.

The p53 gene encodes a 393-amino acid phosphoprotein that can form complexes with host proteins such as large-T antigen and E1B. The protein is found in normal tissues and cells, but at concentrations which are minute by comparison with transformed cells or tumor tissue.

15 Wild-type p53 is recognized as an important growth regulator and a signal transduction protein in many cell types. Missense mutations are common for the p53 gene and are essential for the transforming ability of the oncogene. A single genetic change prompted by point mutations can create carcinogenic p53. Unlike other oncogenes, however, p53 point mutations are known to occur in at least 30
20 distinct codons, often creating dominant alleles that produce shifts in cell phenotype without a reduction to homozygosity. Additionally, many of these dominant negative alleles appear to be tolerated in the organism and passed on in the germ line. Various mutant alleles appear to range from minimally dysfunctional to strongly penetrant, dominant negative alleles (Weinberg, 1991).

25 Another inhibitor of cellular proliferation is p16. The major transitions of the eukaryotic cell cycle are triggered by cyclin-dependent kinases, or CDKs. One CDK, cyclin-dependent kinase 4 (CDK4), regulates progression through G₁. The activity of this enzyme may be to phosphorylate Rb at late G₁. The activity of CDK4 is controlled by an activating subunit, D-type cyclin, and by an inhibitory
30 subunit, the p16^{INK4}, which has been biochemically characterized as a protein that specifically binds to and inhibits CDK4, and thus may regulate retinoblastoma protein (Rb) phosphorylation (Serrano et al., 1993; Serrano et al., 1995). Since the

p16^{INK4} protein is a CDK4 inhibitor (Serrano, 1993), deletion of this gene may increase the activity of CDK4, resulting in hyperphosphorylation of the retinoblastoma protein. p16 also is known to regulate the function of CDK6.

p16^{INK4} belongs to a newly described class of CDK-inhibitory proteins that also includes p16^B, p19, p21^{WAF1}, and p27^{KIP1}. The p16^{INK4} gene maps to 9p21, a chromosome region frequently deleted in many tumor types. Homozygous deletions and mutations of the p16^{INK4} gene are frequent in human tumor cell lines. This evidence suggests that the p16^{INK4} gene is a tumor suppressor gene. This interpretation has been challenged, however, by the observation that the frequency of p16^{INK4} gene alterations is much lower in primary uncultured tumors than in cultured cell lines (Caldas et al., 1994; Cheng et al., 1994; Hussussian et al., 1994; Kamb et al., 1994; Kamb et al., 1994; Mori et al., 1994; Okamoto et al., 1994; Nobori et al., 1995; Orlow et al., 1994; Arap et al., 1995). Restoration of wild-type p16^{INK4} function by transfection with a plasmid expression vector reduced colony formation by some human cancer cell lines (Okamoto, 1994; Arap, 1995).

Other genes that may be employed according to the present invention include Rb, APC, DCC, NF-1, NF-2, WT-1, MEN-I, MEN-II, zac1, p73, VHL, MMAC1 / PTEN, DBCCR-1, FCC, rsk-3, p27, p27/p16 fusions, and p21/p27 fusions.

B.2.f. Regulators of Programmed Cell Death

In certain embodiments, it is contemplated that genetic constructs debilitated for growth, in contrast to wild-type HSV, will stimulate rather than block apoptosis and, thus, will be used to promote the death of diseased or undesired tissue. Apoptosis, or programmed cell death, is an essential process for normal embryonic development, maintaining homeostasis in adult tissues (Kerr et al., 1972). The Bcl-2 family of proteins and the ICE-like proteases have been demonstrated to be important regulators and effectors of apoptosis in other systems. The Bcl-2 protein, discovered in association with follicular lymphoma, plays a prominent role in controlling apoptosis and enhancing cell survival in response to diverse apoptotic stimuli (Bakhshi et al., 1985; Cleary and Sklar, 1985; Cleary et al., 1986; Tsujimoto et al., 1985; Tsujimoto and Croce, 1986). The evolutionarily conserved Bcl-2 protein now is recognized to be a member of a family of related proteins,

which can be categorized as death agonists or death antagonists, *i.e.*, apoptosis regulators.

Subsequent to its discovery, it was shown that Bcl-2 acts to suppress cell death triggered by a variety of stimuli. Also, it now is apparent that there is a family of Bcl-2 cell death regulatory proteins that share common structural and sequence homologies. These different family members have been shown to either possess similar functions to Bcl-2 (*e.g.*, Bcl_{XL}, Bcl_w, Bcl_s, Mcl-1, A1, Bfl-1) or counteract Bcl-2 function and promote cell death (*e.g.*, Bax, Bak, Bik, Bim, Bid, Bad, Harakiri). Wild-type HSV blocks apoptosis by a number of mechanisms, including the modification of proapoptotic members of the Bcl-2 family of proteins.

B.3. PROMOTERS AND ENHANCERS

A "promoter" is a control sequence that is a region of a nucleic acid sequence at which initiation and rate of transcription are controlled. It may contain genetic elements at which regulatory proteins and molecules may bind, such as RNA polymerase and other transcription factors, to initiate the specific transcription a nucleic acid sequence. The phrases "operatively positioned," "operatively linked," "under control," and "under transcriptional control" mean that a promoter is in a correct functional location and/or orientation in relation to a nucleic acid sequence to control transcription of that sequence.

A promoter generally comprises a sequence that functions to position the start site for RNA synthesis. The best known example of this is the TATA box, but in some promoters lacking a TATA box, such as, for example, the promoter for the mammalian terminal deoxynucleotidyl transferase gene and the promoter for the SV40 late genes, a discrete element overlying the start site itself helps to fix the place of initiation. Additional promoter elements regulate the frequency of transcriptional initiation. Typically, these are located in the region 30-110 bp upstream of the start site, although a number of promoters have been shown to contain functional elements downstream of the start site as well. To bring a coding sequence "under the control of" a promoter, one positions the 5' end of the transcription initiation site "downstream" of (*i.e.*, 3') the chosen promoter. The "upstream" promoter stimulates transcription of the DNA and promotes expression of the encoded RNA.

The spacing between promoter elements frequently is flexible, so that promoter function is preserved when elements are inverted or moved relative to one another. In the tk promoter, the spacing between promoter elements can be increased to 50 bp apart before activity begins to decline. Depending on the promoter, it appears that individual elements can function either cooperatively or independently to activate transcription. A promoter may or may not be used in conjunction with an "enhancer," which refers to a cis-acting regulatory sequence involved in the transcriptional activation of a nucleic acid sequence.

A promoter may be one naturally associated with a nucleic acid sequence, as may be obtained by isolating the 5' non-coding sequences located upstream of the coding segment and/or exon. Such a promoter is referred to as "endogenous." Similarly, an enhancer may be one naturally associated with a nucleic acid sequence, located either downstream or upstream of that sequence. Alternatively, certain advantages will be gained by positioning the coding nucleic acid segment under the control of a recombinant or heterologous promoter, which refers to a promoter that is not normally associated with a nucleic acid sequence in its natural environment. A recombinant or heterologous enhancer refers also to an enhancer not normally associated with a nucleic acid sequence in its natural environment. Such promoters or enhancers may include promoters or enhancers of other genes, and promoters or enhancers isolated from any other virus, or prokaryotic or eukaryotic cell, and promoters or enhancers not "naturally occurring," *i.e.*, containing different elements of different transcriptional regulatory regions, consensus sequences, and/or mutations that alter expression. For example, prokaryotic promoters that are most commonly used in recombinant DNA constructions include the β -lactamase (penicillinase), lactose and tryptophan (trp) promoter systems.

Preferred promoters and other expression control elements such as enhancers, transcription factor binding sites, RNA splice sites, polyadenylation signals, terminators, ribosome binding sites and the like (see below), are eukaryotic elements functional in vascular cells. In addition to producing nucleic acid sequences of promoters and enhancers synthetically, sequences may be produced using recombinant cloning and/or nucleic acid amplification technology, including

PCR™, in connection with the compositions disclosed herein (see U.S. Patent Nos. 4,683,202 and 5,928,906, each incorporated herein by reference). Furthermore, it is contemplated that the control sequences that direct transcription and/or expression of sequences within non-nuclear organelles such as mitochondria, chloroplasts, and the like, can be employed as well.

Naturally, it will be important to employ a promoter and/or enhancer that effectively directs the expression of the DNA segment in the organelle, cell type, tissue, organ, or organism chosen for expression. Those of skill in the art of molecular biology generally know the use of promoters, enhancers, and cell type combinations for protein expression (see, for example Sambrook et al. 1989, incorporated herein by reference). The promoters employed may be constitutive, tissue-specific, inducible, and/or useful under the appropriate conditions to direct high level expression of the introduced DNA segment, such as is advantageous in the large-scale production of recombinant proteins and/or peptides. The promoter may be heterologous or endogenous.

In some embodiments, the promoter is constitutive. In other embodiments, the promoter is tissue-specific. In preferred aspects, the promoter is vascular or tissue-specific. The identity of tissue-specific promoters or other elements, as well as assays to characterize their activity, are well known to those of skill in the art. Nonlimiting examples of such elements useful in eukaryotic cells (*e.g.*, mammalian cells) include the SM22 gene promoter (Kuhbandner et al., 2000), human LIMK2 gene (Nomoto et al. 1999), the somatostatin receptor 2 gene (Kraus et al., 1998), murine epididymal retinoic acid-binding gene (Lareyre et al., 1999), human CD4 (Zhao-Emonet et al., 1998), mouse alpha2 (XI) collagen (Tsumaki, et al., 1998), D1A dopamine receptor gene (Lee, et al., 1997), insulin-like growth factor II (Wu et al., 1997), and human platelet endothelial cell adhesion molecule-1 (Almendro et al., 1996).

Additionally, any promoter/enhancer combination (for example, the Eukaryotic Promoter Data Base EPDB, accessed via the Internet) could also be used to drive expression. Use of a T3, T7 or SP6 cytoplasmic expression system is another possible embodiment. Eukaryotic cells can support cytoplasmic transcription from certain bacterial promoters if the appropriate bacterial polymerase

TABLE 4
Promoter and/or Enhancer

Promoter/Enhancer	References
Immunoglobulin Heavy Chain	Banerji <i>et al.</i> , <i>Cell</i> 35:729, 1983; Gilles <i>et al.</i> , 1983; Grosschedl <i>et al.</i> , 1985; Atchinson <i>et al.</i> , 1986, 1987; Imler <i>et al.</i> , 1987; Weinberger <i>et al.</i> , 1984; Kiledjian <i>et al.</i> , 1988; Porton <i>et al.</i> , 1990
Immunoglobulin Light Chain	Queen <i>et al.</i> , 1983; Picard <i>et al.</i> , 1984
T-Cell Receptor	Luria <i>et al.</i> , 1987; Winoto <i>et al.</i> , 1989; Redondo <i>et al.</i> , 1990
HLA DQ a and/or DQ β	Sullivan <i>et al.</i> , 1987
β -Interferon	Goodbourn <i>et al.</i> , 1986; Fujita <i>et al.</i> , 1987; Goodbourn <i>et al.</i> , 1988
Interleukin-2	Greene <i>et al.</i> , 1989
Interleukin-2 Receptor	Greene <i>et al.</i> , 1989; Lin <i>et al.</i> , 1990
MHC Class II 5	Koch <i>et al.</i> , 1989
MHC Class II HLA-Dra	Sherman <i>et al.</i> , 1989
β -Actin	Kawamoto <i>et al.</i> , 1988; Ng <i>et al.</i> , 1989
Muscle Creatine Kinase (MCK)	Jaynes <i>et al.</i> , 1988; Horlick <i>et al.</i> , 1989; Johnson <i>et al.</i> , 1989
Prealbumin (Transthyretin)	Costa <i>et al.</i> , 1988
Elastase I	Omitz <i>et al.</i> , 1987
Metallothionein (MTII)	Karin <i>et al.</i> , 1987; Culotta <i>et al.</i> , 1989
Collagenase	Pinkert <i>et al.</i> , 1987; Angel <i>et al.</i> , <i>Mol. Cell. Biol.</i> 7:2256, 1987]
Albumin	Pinkert <i>et al.</i> , 1987; Tronche <i>et al.</i> , 1989, 1990
α -Fetoprotein	Godbout <i>et al.</i> , 1988; Campere <i>et al.</i> , 1989
γ -Globin	Bodine <i>et al.</i> , 1987; Perez-Stable <i>et al.</i> , 1990
β -Globin	Trudel <i>et al.</i> , 1987
c-fos	Cohen <i>et al.</i> , 1987
c-HA-ras	Triesman, 1986; Deschamps <i>et al.</i> , 1985
Insulin	Edlund <i>et al.</i> , 1985

TABLE 4
Promoter and/or Enhancer

Promoter/Enhancer	References
Neural Cell Adhesion Molecule (NCAM)	Hirsh et al., 1990
α_1 -Antitrypsin	Latimer et al., 1990
H2B (TH2B) Histone	Hwang et al., 1990
Mouse and/or Type I Collagen	Ripe et al., 1989
Glucose-Regulated Proteins (GRP94 and GRP78)	Chang et al., 1989
Rat Growth Hormone	Larsen et al., 1986
Human Serum Amyloid A (SAA)	Edbrooke et al., 1989
Troponin I (TN I)	Yutzey et al., 1989
Platelet-Derived Growth Factor (PDGF)	Pech et al., 1989
Duchenne Muscular Dystrophy	Klamut et al., 1990
SV40	Banerji et al., <i>Cell</i> 27:299, 1981; Moreau et al., 1981; Sleight et al., 1985; Firak et al., 1986; Herr et al., 1986; Imbra et al., 1986; Kadesch et al., 1986; Wang et al., 1986; Ondek et al., 1987; Kuhl et al., 1987; Schaffner et al., 1988
Polyoma	Swartzendruber et al., 1975; Vasseur et al., 1980; Katinka et al., 1980, 1981; Tyndell et al., 1981; Dandolo et al., 1983; de Villiers et al., 1984; Hen et al., 1986; Satake et al., 1988; Campbell and/or Villarreal, 1988
Retroviruses	Kriegler et al., 1982, 1983; Levinson et al., 1982; Kriegler et al., 1983, 1984a, b, 1988; Bosze et al., 1986; Miksicek et al., 1986; Celander et al., 1987; Thiesen et al., 1988; Celander et al., 1988; Chol et al., 1988; Reisman et al., 1989
Papilloma Virus	Campo et al., 1983; Lusky et al., 1983; Spandidos and/or Wilkie, 1983; Spalholz et al., 1985; Lusky et al., 1986; Cripe et al., 1987; Gloss et al., 1987; Hirochika et al., 1987; Stephens et al., 1987; Glue et al., 1988
Hepatitis B Virus	Bulla et al., 1986; Jameel et al., 1986; Shaul et al., 1987; Spandau et al., 1988; Vannice et al., 1988

TABLE 4	
Promoter and/or Enhancer	
Promoter/Enhancer	References
Human Immunodeficiency Virus	Muesing et al., 1987; Hauber et al., 1988; Jakobovits et al., 1988; Feng et al., 1988; Takebe et al., 1988; Rosen et al., 1988; Berkhout <i>et al.</i> , <i>Cell</i> 59:273, 1989; Laspia et al., 1989; Sharp et al., 1989; Braddock et al., 1989
Cytomegalovirus (CMV)	Weber et al., 1984; Boshart et al., 1985; Foecking et al., 1986
Gibbon Ape Leukemia Virus	Holbrook et al., 1987; Quinn et al., 1989

TABLE 5		
Inducible Elements		
Element	Inducer	References
MT II	Phorbol Ester (TFA) Heavy metals	Palmiter et al., 1982; Haslinger et al., 1985; Searle et al., 1985; Stuart et al., 1985; Imagawa et al., 1987, Karin et al., 1987; Angel <i>et al.</i> , <i>Cell</i> 49:729, 1987; McNeall et al., 1989
MMTV (mouse mammary tumor virus)	Glucocorticoids	Huang et al., 1981; Lee et al., 1981; Majors et al., 1983; Chandler et al., 1983; Lee et al., 1984; Ponta et al., 1985; Sakai et al., 1988
β -Interferon	Poly(rI)x Poly(rc)	Tavernier et al., 1983
Adenovirus 5 <u>E2</u>	E1A	Imperiale et al., 1984
Collagenase	Phorbol Ester (TPA)	Angel <i>et al.</i> , <i>Mol. Cell. Biol.</i> 7:2256, 1987
Stromelysin	Phorbol Ester (TPA)	Angel <i>et al.</i> , <i>Cell</i> 49:729, 1987
SV40	Phorbol Ester (TPA)	Angel <i>et al.</i> , <i>Cell</i> 49:729, 1987
Murine MX Gene	Interferon, Newcastle Disease Virus	Hug et al., 1988

TABLE 5
Inducible Elements

Element	Inducer	References
GRP78 Gene	A23187	Resendez et al., 1988
α -2-Macroglobulin	IL-6	Kunz et al., 1989
Vimentin	Serum	Rittling et al., 1989
MHC Class I Gene H-2 κ b	Interferon	Blonar et al., 1989
HSP70	ElA, SV40 Large T Antigen	Taylor et al., 1989, 1990a, 1990b
Proliferin	Phorbol Ester-TPA	Mordacq et al., 1989
Tumor Necrosis Factor α	PMA	Hensel et al., 1989
Thyroid Stimulating Hormone α Gene	Thyroid Hormone	Chatterjee et al., 1989

B.4. INITIATION SIGNALS AND INTERNAL RIBOSOME BINDING SITES

A specific initiation signal also may be required for efficient translation of coding sequences. These signals include the ATG initiation codon or adjacent sequences. Exogenous translational control signals, including the ATG initiation codon, may need to be provided. One of ordinary skill in the art would readily be capable of determining this and providing the necessary signals. It is well known that the initiation codon must be "in-frame" with the reading frame of the desired coding sequence to ensure translation of the entire insert. The exogenous translational control signals and initiation codons can be either natural or synthetic.

In certain embodiments of the invention, the use of internal ribosome entry sites (IRES) elements are used to create multigene, or polycistronic, messages. IRES elements are able to bypass the ribosome scanning model of 5' methylated Cap-dependent translation and begin translation at internal sites (Pelletier and Sonenberg, 1988). IRES elements from two members of the picornavirus family (polio and encephalomyocarditis) have been described (Pelletier and Sonenberg, 1988), as well an IRES from a mammalian message (Macejak and Sarnow, 1991). IRES elements can be linked to heterologous open reading frames. Multiple open reading frames can be transcribed together, each separated by an IRES, creating polycistronic messages. By virtue of the IRES element, each open reading frame is accessible to ribosomes for efficient translation. Multiple genes can be efficiently expressed using a single promoter/enhancer to transcribe a single message (see U.S. Patent Nos. 5,925,565 and 5,935,819, each herein incorporated by reference).

B.5. MULTIPLE CLONING SITES

Vectors can include a multiple cloning site (MCS), which is a nucleic acid region that contains multiple restriction enzyme sites, any of which can be used in conjunction with standard recombinant technology to digest the vector (see, for example, Carbonelli et al., 1999, Levenson et al., 1998, and Cocca, 1997, incorporated herein by reference.) "Restriction enzyme digestion" refers to catalytic cleavage of a nucleic acid molecule with a type II restriction endonuclease that functions only at specific locations in a nucleic acid molecule. Many of these restriction enzymes are commercially available. Use of such enzymes is widely understood by those of skill in the art. Frequently, a vector is linearized or

fragmented using a restriction enzyme that cuts within the MCS to enable exogenous sequences to be ligated to the vector. "Ligation" refers to the process of forming phosphodiester bonds between two nucleic acid fragments. Techniques involving restriction enzymes and ligation reactions are well known to those of skill in the art of recombinant technology.

B.6. SPLICING SITES

Most transcribed eukaryotic RNA molecules will undergo RNA splicing to remove introns from the primary transcripts. Because HSV blocks RNA splicing, cDNAs are the preferred form of eukaryotic insert in the vector, although genomic sequences properly expressible (*e.g.*, intron-free) are also contemplated. Vectors lacking the RNA splice-blocking capacity of wild-type HSV may still require donor and/or acceptor splicing sites to ensure proper processing of the transcript for protein expression (see, for example, Chandler et al., 1997, herein incorporated by reference.)

B.7. TERMINATION SIGNALS

The vectors or constructs of the present invention will generally comprise at least one termination signal. A "termination signal" or "terminator" is comprised of the DNA sequences involved in specific termination of an RNA transcript by an RNA polymerase. Thus, in certain embodiments a termination signal that ends the production of an RNA transcript is contemplated. A terminator may be necessary *in vivo* to achieve desirable message levels.

In eukaryotic systems, the terminator region may also comprise specific DNA sequences that permit site-specific cleavage of the new transcript so as to expose a polyadenylation site. This signals a specialized endogenous polymerase to add a stretch of about 200 A residues (polyA) to the 3' end of the transcript. RNA molecules modified with this polyA tail appear more stable and are translated more efficiently. Thus, in other embodiments involving eukaryotes, it is preferred that that terminator comprises a signal for the cleavage of the RNA, and it is more preferred that the terminator signal promotes polyadenylation of the message. The terminator and/or polyadenylation site elements can serve to enhance message levels and to minimize read-through from a cassette containing a heterologous nucleic acid into other sequences.

Terminators contemplated for use in the invention include any known terminator of transcription described herein or known to one of ordinary skill in the art including, but not limited to, the termination sequences of genes, such as for example the bovine growth hormone terminator or viral termination sequences, such as the SV40 terminator. In certain embodiments, termination may be due to a lack of transcribable or translatable sequence, such as due to a sequence truncation.

B.8. POLYADENYLATION SIGNALS

In expression, particularly eukaryotic expression, one will typically include a polyadenylation signal to effect proper polyadenylation of the transcript. The nature of the polyadenylation signal is not believed to be crucial to the successful practice of the invention, and any such sequence may be employed. Preferred embodiments include the SV40 polyadenylation signal or the bovine growth hormone polyadenylation signal, convenient and known to function well in various target cells. Polyadenylation may increase the stability of the transcript or may facilitate cytoplasmic transport.

B.9. ORIGINS OF REPLICATION

In order to propagate a vector in a host cell, the vector may contain one or more origins of replication sites (often termed "ori"), which is a specific nucleic acid sequence at which replication is initiated. Alternatively, an autonomously replicating sequence (ARS) is employed if the host cell is yeast.

B.10. SELECTABLE AND SCREENABLE MARKERS

In certain embodiments of the invention, cells containing a nucleic acid construct of the present invention may be identified *in vitro* or *in vivo* by including a marker in the expression vector. Such markers confer an identifiable phenotypic change to the cell permitting easy identification of cells containing the expression vector. For example, a screenable marker (*e.g.*, β -galactosidase) may be used to identify transformed cells.

Generally, a selectable marker is one that confers a property that allows for selective survivability. A positive selectable marker is one in which the presence of the marker allows for survival under certain known conditions (*e.g.*, antibiotic challenge), while a negative selectable marker is one in which its presence prevents survival. An example of a positive selectable marker is a drug (*e.g.*, an antibiotic resistance marker).

Usually the inclusion of a drug selection marker aids in the cloning and identification of transformants. For example, genes that confer resistance to neomycin, puromycin, hygromycin, DHFR, GPT, zeocin and histidinol are useful selectable markers. In addition to markers conferring a phenotype that allows for the discrimination of transformants based on the implementation of conditions, other types of markers, including screenable markers such as GFP, whose basis is colorimetric analysis, are also contemplated. Alternatively, screenable enzymes such as herpes simplex virus thymidine kinase (tk) or chloramphenicol acetyltransferase (CAT) may be utilized. One of skill in the art would also know how to employ immunologic markers, possibly in conjunction with FACS analysis. The marker used is not believed to be important, so long as it is capable of being expressed simultaneously with the heterologous nucleic acid encoding a gene product. Further examples of selectable and screenable markers are well known to one of skill in the art.

C. VECTOR DELIVERY AND CELL TRANSFORMATION

Suitable methods for nucleic acid delivery for transformation of an organelle, a cell, a tissue or an organism for use with the current invention are believed to include virtually any method by which a herpes virus vector can be introduced into an organelle, a cell, a tissue or an organism, as described herein or as would be known to one of ordinary skill in the art. Such methods include, but are not limited to, direct delivery of a nucleic acid such as by *ex vivo* transfection (Wilson et al., 1989, Nabel et al, 1989), injection (U.S. Patent Nos. 5,994,624, 5,981,274, 5,945,100, 5,780,448, 5,736,524, 5,702,932, 5,656,610, 5,589,466 and 5,580,859, each incorporated herein by reference), including microinjection (Harlan and Weintraub, 1985; U.S. Patent No. 5,789,215, incorporated herein by reference); by electroporation (U.S. Patent No. 5,384,253, incorporated herein by reference; Turkaspa et al., 1986; Potter et al., 1984); by calcium phosphate precipitation (Graham and Van Der Eb, 1973; Chen and Okayama, 1987; Rippe et al., 1990); by using DEAE-dextran followed by polyethylene glycol (Gopal, 1985); by direct sonic loading (Fechheimer et al., 1987); by liposome-mediated transfection (Nicolau and Sene, 1982; Fraley et al., 1979; Nicolau et al., 1987; Wong et al., 1980; Kaneda et al., 1989; Kato et al., 1991) and receptor-mediated transfection (Wu and Wu, 1987; Wu and Wu, 1988); by microprojectile bombardment (PCT Patent

Application Publication Nos. WO 94/09699 and 95/06128; U.S. Patent Nos. 5,610,042; 5,322,783 5,563,055, 5,550,318, 5,538,877 and 5,538,880, and each incorporated herein by reference), and any combination of such methods. Through the application of techniques such as these, organelle(s), cell(s), tissue(s) or organism(s) may be stably or transiently transformed.

C.1. *EX VIVO* TRANSFORMATION

Methods for transfecting vascular cells and tissues removed from an organism in an *ex vivo* setting are known to those of skill in the art. For example, canine endothelial cells have been genetically altered by retroviral gene transfer *in vitro* and transplanted into a canine (Wilson et al., 1989). In another example, yucatan minipig endothelial cells were transfected by a retrovirus *in vitro* and transplanted into an artery using a double-balloon catheter (Nabel et al., 1989). Thus, it is contemplated that cells or tissues may be removed and transfected *ex vivo* using a herpes virus vector of the present invention. In particular aspects, the transplanted cells or tissues may be placed into an organism. In preferred facets, a therapeutic nucleic acid is expressed in the transplanted cells or tissues.

C.2. INJECTION

In certain embodiments, a nucleic acid may be delivered to an organelle, a cell, a tissue or an organism via one or more injections (*i.e.*, a needle injection), such as, for example, subcutaneously, intradermally, intramuscularly, intravenously, intraperitoneally, and the like. Methods of injection are well known to those of ordinary skill in the art (*e.g.*, injection of a composition comprising a saline solution), and additional examples are described herein.

C.3. LIPOSOME-MEDIATED TRANSFECTION

In a further embodiment of the invention, a nucleic acid may be entrapped in a lipid complex such as, for example, a liposome. Liposomes are vesicular structures characterized by a phospholipid bilayer membrane and an inner aqueous medium. Multilamellar liposomes have multiple lipid layers separated by aqueous medium. They form spontaneously when phospholipids are suspended in an excess of aqueous solution. The lipid components undergo self-rearrangement before the formation of closed structures and entrap water and dissolved solutes between the lipid bilayers (Ghosh and Bachhawat, 1991). Also contemplated is a nucleic acid complexed with Lipofectamine (Gibco BRL) or Superfect (Qiagen).

Liposome-mediated nucleic acid delivery and expression of foreign DNA *in vitro* has been very successful (Nicolau and Sene, 1982; Fraley et al., 1979; Nicolau et al., 1987). The feasibility of liposome-mediated delivery and expression of foreign DNA in cultured chick embryo, HeLa and hepatoma cells has also been demonstrated (Wong et al., 1980).

In certain embodiments of the invention, a liposome may be complexed with a hemagglutinating virus (HVJ). This has been shown to facilitate fusion with the cell membrane and promote cell entry of liposome-encapsulated DNA (Kaneda et al., 1989). In other embodiments, a liposome may be complexed or employed in conjunction with nuclear non-histone chromosomal proteins (HMG-1) (Kato et al., 1991). In yet further embodiments, a liposome may be complexed or employed in conjunction with both HVJ and HMG-1. In other embodiments, a delivery vehicle may comprise a ligand and a liposome. A preferred delivery vehicle comprises a liposome and a ligand selective or, more preferably, specific for a vascular cell-specific binding partner such as a receptor.

C.4. RECEPTOR-MEDIATED TRANSFECTION

Still further, a nucleic acid may be delivered to a target cell via receptor-mediated delivery vehicles. These take advantage of the selective uptake of macromolecules by receptor-mediated endocytosis that will be occurring in a target cell. In view of the cell type-specific distribution of various receptors, this delivery method adds another degree of specificity to the present invention.

Certain receptor-mediated gene-targeting vehicles comprise a cell receptor-specific ligand and a nucleic acid binding agent. Others comprise a cell receptor-specific ligand to which the nucleic acid to be delivered has been operatively attached. Several ligands have been used for receptor-mediated gene transfer (Wu and Wu, 1987; Wagner et al., 1990; Perales et al., 1994; Myers, EPO 0273085), which establishes the operability of the technique. Specific delivery in the context of another mammalian cell type has been described (Wu and Wu, 1993; incorporated herein by reference). In certain aspects of the present invention, a ligand will be chosen to correspond to a receptor specifically expressed on the target cell population.

In other embodiments, a nucleic acid delivery vehicle component of a cell-specific nucleic acid targeting vehicle may comprise a specific binding ligand in

combination with a liposome, as described above. The nucleic acid(s) to be delivered are housed within the liposome and the specific binding ligand is functionally incorporated into the liposome membrane. The liposome will thus specifically bind to the receptor(s) of a target cell and deliver the contents to a cell.

5 Such systems have been shown to be functional using systems in which, for example, epidermal growth factor (EGF) is used in the receptor-mediated delivery of a nucleic acid to cells that exhibit upregulation of the EGF receptor.

In still further embodiments, the nucleic acid delivery vehicle is a liposome itself, which will preferably comprise one or more lipids or glycoproteins that direct cell-specific binding. For example, lactosyl-ceramide, a galactose-terminal

10 asialganglioside, have been incorporated into liposomes and observed to result in an increase in the uptake of the insulin gene by hepatocytes (Nicolau et al., 1987). It is contemplated that the tissue-specific transforming constructs of the present invention are specifically delivered into a target cell in a similar manner.

15 D. HOST CELLS

As used herein, the terms "cell," "cell line," and "cell culture" may be used interchangeably. All of these terms also include their progeny, which is any and all subsequent generations. It is understood that all progeny may not be identical due to deliberate or inadvertent mutations. In the context of expressing a heterologous

20 nucleic acid sequence, "host cell" refers to a prokaryotic or eukaryotic cell (preferred, with vascular cells being particularly preferred), and it includes any transformable organism that is capable of replicating a vector and/or expressing a heterologous nucleic acid encoded by a vector. A host cell can, and has been, used

25 as a recipient for vectors. A host cell may be "infected," "transfected" or "transformed," which refers to a process by which exogenous nucleic acid is transferred or introduced into the host cell. A transformed cell includes the primary subject cell and its progeny. As used herein, the terms "engineered" and "recombinant" cells or host cells are intended to refer to a cell into which an

30 exogenous nucleic acid sequence, such as a vector, has been introduced. Therefore, recombinant cells are distinguishable from naturally occurring cells which do not contain a non-native introduced nucleic acid.

Some vectors may employ control sequences that allow it to be replicated and/or expressed in either or both prokaryotic and eukaryotic cells. One of skill in the art would further understand the conditions under which to incubate all of the above-described host cells to maintain them and to permit replication of a vector (e.g., a herpes virus vector of the invention). Also understood and known are techniques and conditions that would allow large-scale production of vectors, as well as production of the nucleic acids encoded by vectors and their cognate polypeptides, proteins, or peptides.

Numerous cell lines and cultures are available for use as a host cell, and may be obtained through the American Type Culture Collection (ATCC). An appropriate host can be determined by one of skill in the art based on the vector backbone (*i.e.*, the carrier) and the desired result. Examples of eukaryotic host cells for replication and/or expression of a vector include, but are not limited to, HeLa, NIH3T3, Jurkat, 293, Cos, CHO, Saos, and PC12 cells. Many host cells from various cell types and organisms are available and would be known to one of skill in the art.

In certain embodiments, it is contemplated that RNAs or proteinaceous sequences may be co-expressed with other selected RNAs or proteinaceous sequences in the same host cell. Co-expression may be achieved by co-transfecting the host cell with two or more distinct recombinant vectors. Alternatively, a single recombinant vector may be constructed to include multiple distinct coding regions for RNAs, which could then be expressed in host cells transfected with the single vector.

A tissue may comprise a host cell or cells to be transformed with a vector of the present invention. The tissue may be part of, or separated (*e.g.*, by surgery) from, an organism. In certain aspects, cells or a tissue may be removed from a host, contacted with a vector and/or an agent, and then reimplanted into tissue or organism, thereby achieving specificity in vector targeting of host cells through mechanical means such as surgical intervention. In certain embodiments, a tissue may comprise, but is not limited to, adipocytes, alveolar, ameloblasts, axon, basal cells, blood (*e.g.*, lymphocytes), blood vessel, bone, bone marrow, brain, breast, cartilage, cervix, colon, cornea, embryonic, endometrium, endothelial, epithelial, esophagus, facia, fibroblast, follicular, ganglion cells, glial cells, goblet cells,

kidney, liver, lung, lymph node, muscle, neuron, ovaries, pancreas, peripheral blood, prostate, skin, skin, small intestine, spleen, stem cells, stomach, testes cells and all cancers thereof.

The extended duration of expression of a nucleic acid comprised in a herpes virus vector of the present invention in a vascular or a cardiovascular tissue is an advantageous aspect of the present invention. In particular embodiments, a vascular or cardiovascular tissue comprises endothelial cells, smooth muscle cells or adventitial cells. In particular aspects, a vector of the present invention may be expressed in one, two, three or more vascular or cardiovascular cell types, (*e.g.*, endothelial cells, smooth muscle cells or adventitial cells) in any combination of cell types. In certain embodiments, a heterologous nucleic acid sequence may be expressed in a vascular or cardiovascular tissue or cell type for a time greater than or equal to about 1, about 2, about 3, about 4, about 5, about 6, about 7, about 8, about 9, about 10, about 11, about 12, about 13, about 14, about 15, about 16, about 17, about 18, about 19, about 20, about 21, about 22, about 23, about 24, about 25, about 26, about 27, about 28, about 29, about 30, about 31, about 32, about 33, about 34, about 35, about 36, about 37, about 38, about 39, about 40, about 41, about 42, about 43, about 44, about 45, about 46, about 47, about 48, about 49, about 50, about 51, about 52, about 53, about 54, about 55, about 56, about 57, about 58, about 59, about 60, about 61, about 62, about 63, about 64, about 65, about 66, about 67, about 68, about 69, about 70, about 71, about 72, about 73, about 74, about 75, about 76, about 77, about 78, about 79, about 80, about 81, about 82, about 83, about 84, about 85, about 86, about 87, about 88, about 89, about 90, about 91, about 92, about 93, about 94, about 95, about 96, about 97, about 98, about 99, about 100 or more days, and any range derivable therein.

In certain embodiments, the host cell or tissue may be found in at least one organism. In certain embodiments, the organism may be, but is not limited to, an eukaryote (*e.g.*, an animal, a human), as would be understood by one of ordinary skill in the art.

E. THERAPEUTIC AGENTS

In order to increase the effectiveness of a herpes virus vector of the present invention, it may be desirable to combine these compositions and methods of the invention with an agent effective in the treatment of vascular or cardiovascular disease or disorder. In some embodiments, it is contemplated that a conventional therapy or agent including, but not limited to, a pharmacological therapeutic agent, a surgical procedure or a combination thereof, may be combined with vector administration. In a non-limiting example, a therapeutic benefit comprises reduced hypertension in a vascular tissue, or reduced restenosis following vascular or cardiovascular intervention, such as occurs during a medical or surgical procedure. Thus, in certain embodiments, a therapeutic method of the present invention may comprise administration of a vector of the present invention in combination with another therapeutic agent.

This process may involve contacting the cell(s) with an agent(s) and the vector at the same time (*e.g.*, substantially simultaneously) or within a period of time wherein separate administration of the vector and an agent to a cell, tissue or organism produces a desired therapeutic benefit. The terms "contacted" and "exposed," when applied to a cell, tissue or organism, are used herein to describe the process by which a therapeutic construct of the invention and/or therapeutic agent are delivered to a target cell, tissue or organism or are placed in direct contact with the target cell, tissue or organism. The cell, tissue or organism may be contacted (*e.g.*, by administration) with a single composition or pharmacological formulation that includes both a vector of the present invention and one or more agents, or by contacting the cell with two or more distinct compositions or formulations, wherein one composition includes a vector and the other includes one or more agents.

The herpes virus vector may precede, be co-administered with, and/or follow the other agent(s) by intervals ranging from minutes to weeks. In embodiments where the vector and other agent(s) are applied separately to a cell, tissue or organism, one would generally ensure that a significant period of time did not expire between the time of each delivery, such that the vector and agent(s) would still be able to exert an advantageously combined effect on the cell, tissue or organism. For

example, in such instances, it is contemplated that one may contact the cell, tissue or organism with two, three, four or more modalities substantially simultaneously (*i.e.*, within less than about a minute) as the vector. In other aspects, one or more agents may be administered substantially simultaneously with, or about 1 minute, about 5 minutes, about 10 minutes, about 20 minutes about 30 minutes, about 45 minutes, about 60 minutes, about 2 hours, about 3 hours, about 4 hours, about 5 hours, about 6 hours, about 7 hours about 8 hours, about 9 hours, about 10 hours, about 11 hours, about 12 hours, about 13 hours, about 14 hours, about 15 hours, about 16 hours, about 17 hours, about 18 hours, about 19 hours, about 20 hours, about 21 hours, about 22 hours, about 22 hours, about 23 hours, about 24 hours, about 25 hours, about 26 hours, about 27 hours, about 28 hours, about 29 hours, about 30 hours, about 31 hours, about 32 hours, about 33 hours, about 34 hours, about 35 hours, about 36 hours, about 37 hours, about 38 hours, about 39 hours, about 40 hours, about 41 hours, about 42 hours, about 43 hours, about 44 hours, about 45 hours, about 46 hours, about 47 hours, about 48 hours, about 1 day, about 2 days, about 3 days, about 4 days, about 5 days, about 6 days, about 7 days, about 8 days, about 9 days, about 10 days, about 11 days, about 12 days, about 13 days, about 14 days, about 15 days, about 16 days, about 17 days, about 18 days, about 19 days, about 20 days, about 21 days, about 1 week, about 2 weeks, about 3 weeks, about 4 weeks, about 5 weeks, about 6 weeks, about 7 weeks, about 8 weeks, about 1 month, about 2 months, about 3 months, about 4 months, about 5 months, about 6 months, about 7 months, about 8 months, about 9 months, about 10 months, about 11 months, or about 12 months, and any range derivable therein, prior to and/or after administering the vector of the present invention.

Various combination regimens of the herpes virus vectors of the present invention and one or more agents may be employed. Non-limiting examples of such combinations are shown below, wherein a composition comprising a vector is "A" and an agent is "B":

A/B/A	B/A/B	B/B/A	A/A/B	A/B/B	B/A/A	A/B/B/B	B/A/B/B
B/B/B/A	B/B/A/B	A/A/B/B	A/B/A/B	A/B/B/A	B/B/A/A	B/A/B/A	
B/A/A/B	A/A/A/B	B/A/A/A	A/B/A/A	A/A/B/A			

Administration of the composition comprising a vector to a cell, tissue or organism may follow general protocols for the administration of vascular or

cardiovascular therapeutics, taking into account the toxicity of the therapeutic, if any. It is expected that the treatment cycles would be repeated as necessary. It is contemplated that various additional agents may be applied in any combination with the present invention.

5 **E.1. PHARMACOLOGICAL THERAPEUTIC AGENTS**

Pharmacological therapeutic agents and methods of administration, dosages, and the like are well known to those of skill in the art (see for example, the "Physicians Desk Reference," Goodman & Gilman's "The Pharmacological Basis of Therapeutics," "Remington's Pharmaceutical Sciences," and "The Merck Index, 10 Eleventh Edition," incorporated herein by reference in relevant parts), and may be combined with the invention in light of the disclosures herein. Some variation in dosage will necessarily occur depending on the condition of the subject being treated. The person responsible for administration will, in any event, determine the appropriate dose for the individual subject, and such individual determinations are 15 within the skill of those of ordinary skill in the art.

Non-limiting examples of a pharmacological therapeutic agent that may be used in the present invention include an antihyperlipoproteinemic agent, an antiarteriosclerotic agent, an antithrombotic/fibrinolytic agent, a blood coagulant, an antiarrhythmic agent, an antihypertensive agent, a vasopressor, a treatment agent for 20 congestive heart failure, an antianginal agent, an anti-infection agent (*e.g.*, an antibacterial agent, an anti-fungal agent, an anti-viral agent) or a combination thereof.

E.1.a. Antihyperlipoproteinemics

In certain embodiments, administration of an agent that lowers the 25 concentration of one of more blood lipids and/or lipoproteins, known herein as an "antihyperlipoproteinemic," may be combined with administration of a vector for cardiovascular therapy, particularly in treatment of atherosclerosis and thickenings or blockages of vascular tissues. In certain aspects, an antihyperlipoproteinemic agent may comprise an aryloxyalkanoic/fibric acid derivative, a resin/bile acid 30 sequestrant, a HMG CoA reductase inhibitor, a nicotinic acid derivative, a thyroid hormone or thyroid hormone analog, a miscellaneous agent (see below) other antihyperlipoproteinemics known in the art, or a combination thereof.

E.1.a.i. Aryloxyalkanoic Acid/Fibric Acid Derivatives

Non-limiting examples of aryloxyalkanoic/fibric acid derivatives include beclobrate, enzaifibrate, binifibrate, ciprofibrate, clinofibrate, clofibrate (atromide-S), clofibric acid, etofibrate, fenofibrate, gemfibrozil (lobid), nicofibrate, pirifibrate, ronifibrate, simfibrate and theofibrate.

E.1.a.ii. Resins/Bile Acid Sequestrants

Non-limiting examples of resins/bile acid sequestrants include cholestyramine (cholybar, questran), colestipol (colestid) and polidexide.

E.1.a.iii. HMG CoA Reductase Inhibitors

Non-limiting examples of HMG CoA reductase inhibitors include lovastatin (mevacor), pravastatin (pravochol) or simvastatin (zocor).

E.1.a.iv. Nicotinic Acid Derivatives

Non-limiting examples of nicotinic acid derivatives include nicotinate, acepimox, niceritrol, nicoclonate, nicomol and oxiniacic acid.

E.1.a.v. Thyroid Hormones and Analogs

Non-limiting examples of thyroid hormones and analogs thereof include etoroxate, thyropropic acid and thyroxine.

E.1.a.vi. Miscellaneous Antihyperlipoproteinemics

Non-limiting examples of miscellaneous antihyperlipoproteinemics include acifran, azacosterol, benfluorex, β -benzalbutyramide, carnitine, chondroitin sulfate, clomestron, detaxtran, dextran sulfate sodium, 5,8, 11, 14, 17-eicosapentaenoic acid, eritadenine, furazabol, meglutol, melinamide, mytatrienediol, ornithine, γ -oryzanol, pantethine, pentaerythritol tetraacetate, α -phenylbutyramide, pirozadil, probucol (lorelco), β -sitosterol, sultosilic acid-piperazine salt, tiadenol, triparanol and xenbucin.

E.1.b. Antiarteriosclerotics

A non-limiting example of an antiarteriosclerotic is pyridinol carbamate.

E.1.c. Antithrombotic/Fibrinolytic Agents

In certain embodiments, administration of an agent that aids in the removal or prevention of blood clots may be combined with administration of a vector for cardiovascular therapy, particularly in treatment of atherosclerosis and vasculature (e.g., arterial) blockages. Non-limiting examples of antithrombotic and/or

fibrinolytic agents include anticoagulants, anticoagulant antagonists, antiplatelet agents, thrombolytic agents, thrombolytic agent antagonists or combinations thereof.

In certain aspects, antithrombotic agents that can be administered orally, such as, aspirin and warfarin (coumadin), are preferred.

5 **E.1.c.i.. Anticoagulants**

Non-limiting examples of an anticoagulant include acenocoumarol, ancrod, anisindione, bromindione, clorindione, coumetarol, cyclocoumarol, dextran sulfate sodium, dicoumarol, diphenadione, ethyl biscoumacetate, ethylidene dicoumarol, fluidione, heparin, hirudin, lyapolate sodium, oxazidione, pentosan polysulfate, 10 phenindione, phenprocoumon, phosvitin, picotamide, tiocloamarol and warfarin.

E.1.c.ii. Antiplatelet Agents

Non-limiting examples of antiplatelet agents include aspirin, a dextran, dipyridamole (persantin), heparin, sulfinpyranone (anturane) and ticlopidine (ticlid).

E.1.c.iii. Thrombolytic Agents

15 Non-limiting examples of thrombolytic agents include tissue plasminogen activator (activase), plasmin, pro-urokinase, urokinase (abbokinase), streptokinase (streptase), and anistreplase/APSAC (eminase).

E.1.d. Blood Coagulants

In certain embodiments wherein a patient is suffering from a hemorrhage or 20 an increased likelihood of hemorrhaging, an agent that may enhance blood coagulation may be used. Non-limiting examples of a blood coagulation-promoting agent include thrombolytic agent antagonists and anticoagulant antagonists.

E.1.d.i. Anticoagulant Antagonists

Non-limiting examples of anticoagulant antagonists include protamine and 25 vitamin K1.

E.1.d.ii. Thrombolytic Agent Antagonists and Antithrombotics

Non-limiting examples of thrombolytic agent antagonists include aminocaproic acid (amicar) and transexamic acid (amstat). Non-limiting examples of antithrombotics include anagrelide, argatroban, cilostazol, daltroban, defibrotide, 30 enoxaparin, fraxiparine, indobufen, lamoparan, ozagrel, picotamide, plafibride, tedelparin, ticlopidine and triflusal.

E.1.e. Antiarrhythmic Agents

Non-limiting examples of antiarrhythmic agents include Class I antiarrhythmic agents (sodium channel blockers), Class II antiarrhythmic agents (beta-adrenergic blockers), Class III antiarrhythmic agents (repolarization prolonging drugs), Class IV antiarrhythmic agents (calcium channel blockers) and miscellaneous antiarrhythmic agents (see below).

E.1.e.i. Sodium Channel Blockers

Non-limiting examples of sodium channel blockers include Class IA, Class IB and Class IC antiarrhythmic agents. Non-limiting examples of Class IA antiarrhythmic agents include disopyramide (norpace), procainamide (pronestyl) and quinidine (quinidex). Non-limiting examples of Class IB antiarrhythmic agents include lidocaine (xylocaine), tocainide (tonocard) and mexiletine (mexitil). Non-limiting examples of Class IC antiarrhythmic agents include encainide (enkaid) and flecainide (tambocor).

E.1.e.ii. Beta Blockers

Non-limiting examples of a beta blocker, otherwise known as a β -adrenergic blocker, a β -adrenergic antagonist or a Class II antiarrhythmic agent, include acebutolol (sectral), alprenolol, amosulalol, arotinolol, atenolol, befunolol, betaxolol, bevantolol, bisoprolol, bopindolol, bucumolol, bufetolol, bufuralol, bunitrolol, bupranolol, butidine hydrochloride, butofilolol, carazolol, carteolol, carvedilol, celiprolol, cetamolol, cloranolol, dilevalol, epanolol, esmolol (brevibloc), indenolol, labetalol, levobunolol, mepindolol, metipranolol, metoprolol, moprolool, nadolol, nadoxolol, nifenalol, nipradilol, oxprenolol, penbutolol, pindolol, practolol, pronethalol, propanolol (inalderal), sotalol (betapace), sulfinalol, talinolol, tertatolol, timolol, toliprolol and xibinolol. In certain aspects, the beta blocker comprises an aryloxypropanolamine derivative. Non-limiting examples of aryloxypropanolamine derivatives include acebutolol, alprenolol, arotinolol, atenolol, betaxolol, bevantolol, bisoprolol, bopindolol, bunitrolol, butofilolol, carazolol, carteolol, carvedilol, celiprolol, cetamolol, epanolol, indenolol, mepindolol, metipranolol, metoprolol, moprolool, nadolol, nipradilol, oxprenolol, penbutolol, pindolol, propanolol, talinolol, tertatolol, timolol and toliprolol.

E.1.e.iii. Repolarization Prolonging Agents

Non-limiting examples of an agent that prolong repolarization, also known as a Class III antiarrhythmic agent, include amiodarone (cordarone) and sotalol (betapace).

5 E.1.e.iv. Calcium Channel Blockers/Antagonist

Non-limiting examples of a calcium channel blocker, otherwise known as a Class IV antiarrhythmic agent, include an arylalkylamine (*e.g.*, bepridile, diltiazem, fendiline, gallopamil, prenylamine, terodiline, verapamil), a dihydropyridine derivative (felodipine, isradipine, nicardipine, nifedipine, nimodipine, nisoldipine, nitrendipine) a piperazine derivative (*e.g.*, cinnarizine, flunarizine, lidoflazine) or a miscellaneous calcium channel blocker such as bencyclane, etafenone, magnesium, mibefradil or perhexiline. In certain embodiments a calcium channel blocker comprises a long-acting dihydropyridine (nifedipine-type) calcium antagonist.

E.1.e.v. Miscellaneous Antiarrhythmic Agents

15 Non-limiting examples of miscellaneous antiarrhythmic agents include adenosine (adenocard), digoxin (lanoxin), acecainide, ajmaline, amoproxan, aprindine, bretylium tosylate, bunaftine, butobendine, capobenic acid, cifenline, disopyranide, hydroquinidine, indecainide, ipatropium bromide, lidocaine, lorajmine, lorcainide, meobentine, moricizine, pirmenol, prajmaline, propafenone, pyrinoline, quinidine polygalacturonate, quinidine sulfate and viquidil.

E.1.f. Antihypertensive Agents

Non-limiting examples of antihypertensive agents include sympatholytic, alpha/beta blockers, alpha blockers, anti-angiotensin II agents, beta blockers, calcium channel blockers, vasodilators and miscellaneous antihypertensives.

25 E.1.f.i. Alpha Blockers

Non-limiting examples of an alpha blocker, also known as an α -adrenergic blocker or an α -adrenergic antagonist, include amosulalol, arotinolol, dapiprazole, doxazosin, ergoloid mesylates, fenspiride, indoramin, labetalol, nicergoline, prazosin, terazosin, tolazoline, trimazosin and yohimbine. In certain embodiments, an alpha blocker may comprise a quinazoline derivative. Non-limiting examples of quinazoline derivatives include alfuzosin, bunazosin, doxazosin, prazosin, terazosin and trimazosin.

E.1.f.ii. Alpha/Beta Blockers

In certain embodiments, an antihypertensive agent is both an alpha and beta adrenergic antagonist. A non-limiting example of an alpha/beta blocker is labetalol (normodyne, trandate).

5 E.1.f.iii. Anti-Angiotension II Agents

Non-limiting examples of anti-angiotension II agents include angiotensin converting enzyme inhibitors and angiotension II receptor antagonists. Non-limiting examples of angiotension converting enzyme inhibitors (ACE inhibitors) include alacepril, enalapril (vasotec), captopril, cilazapril, delapril, enalaprilat, fosinopril, 10 lisinopril, moveltopril, perindopril, quinapril and ramipril. Non-limiting examples of an angiotensin II receptor blocker, also known as an angiotension II receptor antagonist, an ANG receptor blocker or an ANG-II type-1 receptor blocker (ARBS), include angiocandesartan, eprosartan, irbesartan, losartan and valsartan.

E.1.f.iv. Sympatholytics

15 Non-limiting examples of a sympatholytic include a centrally acting sympatholytic or a peripherally acting sympatholytic. Non-limiting examples of a centrally acting sympatholytic, also known as a central nervous system (CNS) sympatholytic, include clonidine (catapres), guanabenz (wytensin) guanfacine (tenex) and methyldopa (aldomet). Non-limiting examples of a peripherally acting 20 sympatholytic include a ganglion-blocking agent, an adrenergic neuron-blocking agent, a β -adrenergic blocking agent or an α 1-adrenergic blocking agent. Non-limiting examples of a ganglion-blocking agent include mecamylamine (inversine) and trimethaphan (arfonad). Non-limiting examples of an adrenergic neuron-blocking agent include guanethidine (ismelin) and reserpine (serpasil). Non-limiting 25 examples of a β -adrenergic blocker include acenitolol (sectral), atenolol (tenormin), betaxolol (kerlone), carteolol (cartrol), labetalol (normodyne, trandate), metoprolol (lopressor), nadanol (corgard), penbutolol (levatol), pindolol (visken), propranolol (inderal) and timolol (blocadren). Non-limiting examples of α 1-adrenergic blockers include prazosin (minipress), doxazocin (cardura) and terazosin (hytrin).

30 E.1.f.v. Vasodilators

In certain embodiments, a cardiovascular therapeutic agent may comprise a vasodilator (e.g., a cerebral vasodilator, a coronary vasodilator or a peripheral vasodilator). In certain preferred embodiments, a vasodilator comprises a coronary

vasodilator. Non-limiting examples of a coronary vasodilator include amotriphene, bendazol, benfurodil hemisuccinate, benziodarone, chloracizine, chromonar, clobenfurol, clonitrate, dilazep, dipyridamole, droprenilamine, efloxate, erythrityl tetranitrane, etafenone, fendiline, floredil, ganglefene, herestrol bis(β -diethylaminoethyl ether), hexobendine, itramin tosylate, khellin, lidoflanine, mannitol hexanitrane, medibazine, nicorglycerin, pentaerythritol tetranitrate, pentrinitrol, perhexiline, pimefylline, trapidil, tricromyl, trimetazidine, trolnitrate phosphate and visnadine.

In certain aspects, a vasodilator may comprise a chronic therapy vasodilator or a hypertensive emergency vasodilator. Non-limiting examples of a chronic therapy vasodilator include hydralazine (apresoline) and minoxidil (loniten). Non-limiting examples of a hypertensive emergency vasodilator include nitroprusside (nipride), diazoxide (hyperstat IV), hydralazine (apresoline), minoxidil (loniten) and verapamil.

E.1.f.vi. Miscellaneous Antihypertensives

Non-limiting examples of miscellaneous antihypertensives include ajmaline, γ -aminobutyric acid, bufeniode, cicletanine, ciclosidomine, a cryptenamine tannate, fenoldopam, flosequinan, ketanserin, mebutamate, mecamlamine, methyldopa, methyl 4-pyridyl ketone thiosemicarbazone, muzolimine, pargyline, pempidine, pinacidil, piperoxan, primaperone, a protoveratrine, raubasine, rescimetol, rilmenidene, saralasin, sodium nitroprusside, ticrynafen, trimethaphan camsylate, tyrosinase and urapidil.

In certain aspects, an antihypertensive may comprise an arylethanolamine derivative (see below), a benzothiadiazine derivative, a N-carboxyalkyl(peptide/lactam) derivative, a dihydropyridine derivative, a guanidine derivative, a hydrazines/phthalazine, an imidazole derivative, a quaternary ammonium compound, a reserpine derivative or a sulfonamide derivative.

E.1.f.vi.a. Arylethanolamine Derivatives

Non-limiting examples of arylethanolamine derivatives include amosulalol, bufuralol, dilevalol, labetalol, pronethalol, sotalol and sulfinalol.

E.1.f.vi.b. Benzothiadiazine Derivatives

Non-limiting examples of benzothiadiazine derivatives include althizide, bendroflumethiazide, benzthiazide, benzyhydrochlorothiazide, buthiazide,

chlorothiazide, chlorthalidone, cyclopenthiiazide, cyclothiazide, diazoxide, epithiazide, ethiazide, fenquizone, hydrochlorothizide, hydroflumethizide, methyclothiazide, meticrane, metolazone, paraflutizide, polythizide, tetrachlormethiazide and trichlormethiazide.

5 **E..1.f.vi.c. N-carboxyalkyl(peptide/lactam) Derivatives**

Non-limiting examples of N-carboxyalkyl(peptide/lactam) derivatives include alacepril, captopril, cilazapril, delapril, enalapril, enalaprilat, fosinopril, lisinopril, moveltipril, perindopril, quinapril and ramipril.

E.1.f.vi.d. Dihydropyridine Derivatives

10 Non-limiting examples of dihydropyridine derivatives include amlodipine, felodipine, isradipine, nicardipine, nifedipine, nilvadipine, nisoldipine and nitrendipine.

E.1.f.vi.e. Guanidine Derivatives

15 Non-limiting examples of guanidine derivatives include bethanidine, debrisoquin, guanabenz, guanacine, guanadrel, guanazodine, guanethidine, guanfacine, guanochlor, guanoxabenz and guanoxan.

E.1.f.vi.f. Hydrazines/Phthalazines

20 Non-limiting examples of hydrazines/phthalazines include budralazine, cadralazine, dihydralazine, endralazine, hydracarbazine, hydralazine, pheniprazine, pildralazine and todralazine.

E.1.f.vi.g. Imidazole Derivatives

Non-limiting examples of imidazole derivatives include clonidine, lofexidine, phentolamine, tiamenidine and tolonidine.

E.1.f.vi.h. Quaternary Ammonium Compounds

25 Non-limiting examples of quaternary ammonium compounds include azamethonium bromide, chlorisondamine chloride, hexamethonium, pentacynium bis(methylsulfate), pentamethonium bromide, pentolinium tartrate, phenactropinium chloride and trimethidinium methosulfate.

E.1.f.vi.i. Reserpine Derivatives

30 Non-limiting examples of reserpine derivatives include bietaserpine, deserpidine, rescinnamine, reserpine and syrosingopine.

E.1.f.vi.j. Sulfonamide Derivatives

Non-limiting examples of sulfonamide derivatives include ambuside, clopamide, furosemide, indapamide, quinethazone, tripamide and xipamide.

E.1.g. Vasopressors

5 Vasopressors generally are used to increase blood pressure during shock, which may occur during a surgical procedure. Non-limiting examples of a vasopressor, also known as an antihypotensive, include amezinium methyl sulfate, angiotensin amide, dimetofrine, dopamine, etifelmin, etilefrin, gepefrine, metaraminol, midodrine, norepinephrine, pholedrine and synephrine.

E.1.h. Treatment Agents for Congestive Heart Failure

Non-limiting examples of agents for the treatment of congestive heart failure include anti-angiotension II agents, afterload-preload reduction treatment (see below), diuretics and inotropic agents.

E.1.h.i. Afterload-Preload Reduction Treatment

15 In certain embodiments, an animal (patient) that cannot tolerate an angiotension antagonist may be treated with a combination therapy. Such therapy may combine administration of hydralazine (apresoline) and isosorbide dinitrate (isordil, sorbitrate).

E.1.h.ii. Diuretics

20 Non-limiting examples of a diuretic include a thiazide or benzothiadiazine derivative (e.g., althiazide, bendroflumethazide, benzthiazide, benzyhydrochlorothiazide, buthiazide, chlorothiazide, chlorothiazide, chlorthalidone, cyclopenthiazide, epithiazide, ethiazide, ethiazide, fenquizone, hydrochlorothiazide, hydroflumethiazide, methyclothiazide, meticrane, metolazone, paraflutizide, polythizide, tetrachloromethiazide, trichlormethiazide), an
25 organomercurial (e.g., chlormerodrin, meralluride, mercamphamide, mercaptomerin sodium, mercumallylic acid, mercumatilin sodium, mercurous chloride, mersalyl), a pteridine (e.g., furterene, triamterene), purines (e.g., acefylline, 7-morpholinomethyltheophylline, pamobrom, protheobromine, theobromine),
30 steroids including aldosterone antagonists (e.g., canrenone, oleandrin, spironolactone), a sulfonamide derivative (e.g., acetazolamide, ambuside, azosemide, bumetanide, butazolamide, chloraminophenamide, clofenamide, clopamide, clorexolone, diphenylmethane-4,4'-disulfonamide, disulfamide,

ethoxazolamide, furosemide, indapamide, mefruside, methazolamide, piretanide, quinethazone, torasemide, triamide, xipamide), a uracil (*e.g.*, aminometradine, amisometradine), a potassium-sparing antagonist (*e.g.*, amiloride, triamterene) or a miscellaneous diuretic such as aminozine, arbutin, chlorazasil, ethacrynic acid, etozolin, hydracarbazine, isosorbide, mannitol, metochalcone, muzolimine, perhexiline, ticnafen and urea.

E.1.h.iii. Intropic Agents

Non-limiting examples of a positive inotropic agent, also known as a cardiotonic, include acefylline, an acetyldigitoxin, 2-amino-4-picoline, amrinone, benfurodil hemisuccinate, bucladesine, cerberosine, camphotamide, convallatoxin, cymarin, denopamine, deslanoside, digitalin, digitalis, digitoxin, digoxin, dobutamine, dopamine, dopexamine, enoximone, erythrophleine, fenalcomine, gitalin, gitoxin, glycyamine, heptaminol, hydrastinine, ibopamine, a lanatoside, metamivam, milrinone, nerifolin, oleandrin, ouabain, oxyfedrine, prenalterol, proscillaridine, resibufogenin, scillaren, scillarenin, strphanthin, sulmazole, theobromine and xamoterol.

In particular aspects, an inotropic agent is a cardiac glycoside, a beta-adrenergic agonist or a phosphodiesterase inhibitor. Non-limiting examples of a cardiac glycoside includes digoxin (lanoxin) and digitoxin (crystodigin). Non-limiting examples of a β -adrenergic agonist include albuterol, bambuterol, bitolterol, carbuterol, clenbuterol, clorprenaline, denopamine, dioxethedrine, dobutamine (dobutrex), dopamine (intropin), dopexamine, ephedrine, etafedrine, ethylnorepinephrine, fenoterol, formoterol, hexoprenaline, ibopamine, isoetharine, isoproterenol, mabuterol, metaproterenol, methoxyphenamine, oxyfedrine, pirbuterol, procaterol, protokylol, reproterol, rimiterol, ritodrine, soterenol, terbutaline, tretoquinol, tulobuterol and xamoterol. A non-limiting example of a phosphodiesterase inhibitor is amrinone (inocor).

E.1.i. Antianginal Agents

Antianginal agents may comprise organonitrates, calcium channel blockers, beta blockers and combinations thereof.

Non-limiting examples of organonitrates, also known as nitrovasodilators, include nitroglycerin (nitro-bid, nitrostat), isosorbide dinitrate (isordil, sorbitrate) and amyl nitrate (aspirol, vaporole).

E.1.j. Anti-Infection Agents

Anti-infection agents (*e.g.*, antibacterials, antifungus, antivirals) are generally used to reduce or prevent infection. Non-limiting examples of antibacterials include antibacterial antibiotics, synthetic antibacterials, leprostatic
5 antibacterials, rickettsia antibacterials, tuberculostatic antibacterials, or a combination thereof.

E.1.j.i. Antibiotic Antibacterials

Non-limiting examples of antibiotic antibacterials include an aminoglycoside (*e.g.*, amikacin, apramycin, arbekacin, a bambermycin, butirosin, dibekacin,
10 dihydrostreptomycin, a fortimicin, gentamicin, isepamicin, kanamycin, micromycin, neomycin undecylenate, netilmicin, paromomycin, ribostamycin, sisomicin, spectinomycin, streptomycin, streptonicozid, tobramycin), an amphenol (*e.g.*, azidamfenicol, chloramphenicol, chloramphenicol palmitate, chloramphenicol pantothenate, florfenicol, thiamphenicol), an ansamycin (*e.g.*, rifamide, rifampin,
15 rifamycin, rifaximin), a β -lactam (*e.g.*, a carbapenem, a cephalosporin, a cephamycin, a monobactam, an oxacephem, a penicillin), a lincosamide (*e.g.*, clindamycin, lincomycin), a macrolide (*e.g.*, azithromycin, carbomycin, clarithromycin, erythromycin acistrate, erythromycin estolate, erythromycin glucoheptonate, erythromycin lactobionate, erythromycin lactobionate, erythromycin
20 propionate, erythromycin stearate, josamycin, leucomycin, midecamycin, miokamycin, oleandomycin, primycin, primycin, rokitamycin, rosaramicin, roxithromycin, spiramycin, troleandomycin), polypeptides (*e.g.*, amphomycin, bacitracin, capreomycin, colistin, enduracidin, enviomycin, fusagungine, a gramicidin, a gramicidin S, mikamycin, polymyxin, polymyxin B-Methanesulfonic
25 acid, pristnamycin, ristocetin, teicoplanin, thiostrepton, tuberactinomycin, tyrocidine, tyrothricin, vancomycin, viomycin, viomycin pantothenate, virginiamycin, zinc bacitracin), tetracycline (*e.g.*, apicycline, chlortetracycline, clomocycline, demeclocycline, doxycycline, guamecycline, lymecycline, meclocycline, methacycline, minocycline, oxytetracycline, penimepicycline,
30 pipacycline, rolitetracycline, sancycline, senociclin, tetracycline) or a miscellaneous antibiotic antibacterial (*e.g.*, cycloserin, mupirocin, tuberin).

A non-limiting example of a carbapenem β -lactam is imipenem. Non-limiting examples of a cephalosporin β -lactam include cefaclor, cefadroxil,

cefamandole, cefatrizine, cefazedone, cefazolin, cefixime, cefmenoxime, cefodizime, cefonicid, cefoperazone, ceforanide, cefotaxime, cefotiam, cefpimizole, cefpiramide, cefpodoxime proxetil, cefroxadine, cefsulodin, ceftazidime, cefteram, cftezole, ceftibuten, ceftizoxime, ceftriaxone, cefuroxime, cefuzonam, cephaetrile sodium, cephalixin, cephaloglycin, cephaloridine, cephalosporin C, cephalothin, cephapirin sodium, cephradine and pivcefalexin. Non-limiting examples of a cephamycin β -lactam include cefbuperazone, cefmetazole, cefminox, cefotetan and cefoxitin. Non-limiting examples of a monobactam β -lactam include aztreonam, carumonam and tigemonam. Non-limiting examples of an oxacephem β -lactam include flomoxef and moxolactam. Non-limiting examples of a penicillin β -lactam include amidinocillin, amidinocillin pivoxil, amoxicillin, ampicillin, apalcillin, aspoxicillin, azidocillin, azlocillin, bacampicillin, benzylpenicillinic acid, benzylpenicillin sodium, carbenicillin, carfecillin sodium, carindacillin, clometocillin, cloxacillin, cyclacillin, dicloxacillin diphenicillin sodium, epicillin, fenbenicillin, floxacillin, hetacillin, lenampicillin, metampicillin, methicillin sodium, mezlocillin, nafcillin sodium, mezlocillin, nafcillin sodium, oxacillin, penamecillin, penethamate hydride, penicillin G benethiamine, penicillin G benzathine, penicillin G benzhydrylamine, penicillin G calcium, penicillin G hydrabamine, penicillin G potassium, penicillin G procaine, penicillin N, penicillin O, penicillin V, penicillin V benzathine, penicillin V hydrabamine, penimepicycline, phenethicillin potassium, piperacillin, pivampicillin, propicillin, quinacillin, sulbenicillin, talampicillin, temocillin and ticarcillin.

E.1.j.ii. Synthetic Antibacterials

Non-limiting examples of synthetic antibacterials include 2,4-diaminopyrimidines (*e.g.*, brodimoprim, tetroxoprim, trimethoprim), nitrofurans (*e.g.*, furaltadone, furazolium chloride, nifuradene, nifuratel, nifurfoline, nifurpirinol, nifurpazine, nifurtoinol, nitrofurantion), quinolones and quinone analogs (*e.g.*, amifoxacin, cinoxacin, ciprofloxacin, difloxacin, enoxacin, fleroxacin, flumequine, lomefloxacin, miloxacin, nalidixic acid, norfloxacin, ofloxacin, oxolinic acid, pefloxacin, pipemidic acid, piromidic acid, rosoxacin, temafloxacin, tosulfoxacin), sulfonamides (*e.g.*, acetyl sulfamehtoxypraxine, acetyl sulfisoxazole, azosulfamide, benzylsulfamide, choramine-B, chloramine-T, dichloramine T, formosulfathiazole, N²-formylsulfisomidine, N⁴- β -D-

glucosylsulfanilamide, mafenide, 4'-(methylsulfanoyl)sulfanilamide, p-nitrosulfathiazole, phthalylsulfacetamide, phthalylsulfathiazole, salazosulfadimidine, succinylsulfathiazole, sulfabenzamide, sulfacetamide, sulfachlorpyridazine, sulfachrysoidine, sulfacytine, sulfadiazine, sulfadicramide, sulfadimethoxine, sulfadoxine, sulfaethidole, sulfaguanidine, sulfaguanol, sulfalene, sulfaloxic acid, sulfamerazine, sulfameter, sulfamethazine, sulfamethizole, sulfamethomidine, sulfamethoxazole, sulfamethoxypyridazine, sulfametrole, sulfamidochrysoidine, sulfamoxole, sulfanilamide, sulfanilamidomethanesulfonic acid triethanolamine salt, 4-sulfanilamidosalicylic acid, N⁴-sulfanilylsulfanilamide, sulfanilylurea, N-sulfanilyl-3,4-xylamide, sulfanitran, sulfaperine, sulfaphenazole, sulfaproxyline, sulfapyrazine, sulfapyridine, sulfasomizole, sulfasymazine, sulfathiazole, sulfathiourea, sulfatolamide, sulfisomidine, sulfisoxazole), sulfones (acedapsone, acediasulfone, acetosulfone sodium, dapsone, diathymosulfone, glucosulfone sodium, solasulfone, succisulfone, sulfanilic acid, p-sulfanilylbenzylamine, p,p'-sulfonyldianiline-N,N'-diagalactoside, sulfoxone sodium, thiazolsulfone), and miscellaneous synthetic antibacterials (*e.g.*, clofoctol, hexedine, methenamine, methenamine anhydromethylene-citrate, methenamine hippurate, methenamine mandelate, methenamine sulfosalicylate, nitroxoline, and xibornol).

E.1.j.iii. Liprostatic Antibacterials

Non-limiting examples of leprostatic antibacterials include acedapsone, acetosulfone sodium, clofazimine, dapsone, diathymosulfone, glucosulfone sodium, hydnocarpic acid, solasulfone, succisulfone and sulfoxone sodium.

E.1.j.iv. Rickettsia Antibacterials

Non-limiting examples of rickettsia antibacterials, also known as antirickettsials, include p-aminobenzoic acid, chloramphenicol, chloramphenicol palmitate, chloramphenicol pantothenate and tetracycline.

E.1.j.v. Tuberculostatic Antibacterials

Non-limiting examples of tuberculostatic antibacterials include p-aminosalicylic acid, p-aminosalicylic acid hydrazine, benzoylpas, 5-bromosalicylhydroxamic acid, capreomycin, clofazimine, cyacetacide, cycloserine, dihydrostreptomycin, enviomycin, ethambutol, ethionamide, 4'-formylsuccinanilic acid thiosemicarbazone, furonazide, glyconiazide, isobutol, isoniazide, isoniazid methanesulfonate, morphazinamide, opiniazide, parsiniazide, phenyl

aminosalicylate, protionamide, pyrazinamide, rifampin, salinazide, streptomycin, subathizone, sulfoniazide, thiacetazone, tiocarlide, tuberactinomycin, tubercidin, tuberin verazide, viomycin and vicmycin pantothenate.

E.1.j.vi. Antifungal Agents

5 Examples of antifungal agents include antibiotic antifungal agents and synthetic antifungal agents. Non-limiting examples of antibiotic antifungal agents include polyenes, such as amphotericin B, candicidin, dermostatin, filipin, fungichromin, hachimycin, hamycin, lucensomycin, mepartricin, natamycin, nystatin, pecilocin and perimycin; as well as miscellaneous antibiotic antifungal agents such as azaserine, grisofulvin, an oligomycin, neomycin undecylenate, 10 pyrrolnitrin, siccanin, tubercidin and viridin. Examples of synthetic antifungal agents include allylamines, imidazoles, triazoles and miscellaneous synthetic antifungal agents. Non-limiting examples of allylamines include naftifine and terbinafine. Non-limiting examples of imidazoles include bifonazole, butoconazole, 15 chlordantoin, chlormidazole, cloconazole, clotrimazole, econazole, enilconazole, fenticonazole, isoconazole, ketoconazole, miconazole, omoconazole, oxiconazole nitrate, sulconazole and tioconazole. Non-limiting examples of trizoles include fluconazole, itraconazole and terconazole. Non-limiting examples of miscellaneous synthetic antifungal agents include acrisorcin, amorolfine, biphenamine, 20 bromosalicylchloranilide, buclosamide, calcium propionate, chlorphenesin, ciclopirox, cloxyquin, coparaffinate, diamthazole dihydrochloride, exalamide, flucytosine, halethazole, hexetidine, loflucarban, nifuratel, potassium iodide, propionic acid, pyrrithione, salicylanilide, sodium propionate, sulbentine, tenonitrozole, tolciolate, tolindate, tolnaftate, triacetin, ujothion, undecylenic acid 25 and zinc propionate.

E.1.j.vii. Antiviral Agents

 Non-limiting example of antiviral agents include certain purines and pyrimidinones, such as acyclovir, cytarabine, dideoxyadenosine, dideoxycytidine, dideoxyinosine, edoxudine, floxuridine, ganciclovir, idoxuridine, inosine pranobex, 30 MADU, trifluridine, vidarabine and zidovudine. Miscellaneous antiviral agents include acetylleucine monoethanolamine, amantadine, amidinomycin, cuminaldehyde, thiosemicarbazone, foscarnet sodium, interferon- α , interferon- β ,

interferon- γ , kethoxal, lysozyme, methisazone, moroxydine, podophyllotoxin, ribavirin, rimantadine, stallimycin, statolon, tromantadine and xenazoic acid.

E.2. Surgical Procedures

5 In certain aspects, surgical procedures may be performed along with the administration of the HSV vector, or to facilitate targeting of the vector to vascular cells, tissues and organs. Surgical procedures would include, for example, preventative, diagnostic or staging, curative, and palliative surgery. Surgery, and in particular a curative surgery, may be used in conjunction with other therapies, such as administraction of a growth-debilitated (CNS) HSV vector and one or more other
10 agents.

Such surgical procedures for vascular and cardiovascular diseases and disorders are well known to those of skill in the art, and include, but are not limited to, performing surgery on an organism, providing a cardiovascular mechanical prosthesis, angioplasty, coronary artery reperfusion, catheter ablation, providing an
15 implantable cardioverter defibrillator to the subject, mechanical circulatory support, or a combination thereof. Non-limiting examples of a mechanical circulatory support that may be used in the present invention comprise an intra-aortic balloon counterpulsation, left ventricular assist device or combination thereof.

Of course, further treatment of the area of surgery may be accomplished by
20 perfusion, direct injection, systemic injection or local application of the area with at least one additional therapeutic agent (*e.g.*, a vector for use in the methods of the invention, a pharmacological therapeutic agent), as would be known to one of skill in the art or described herein.

F. PHARMACEUTICAL PREPARATIONS

25 Pharmaceutical compositions of the present invention comprise an effective amount of one or more herpes virus vector(s) or additional agent(s) dissolved or dispersed in a pharmaceutically acceptable carrier. The phrases "pharmaceutical or pharmacologically acceptable" refer to molecular entities and compositions that do
30 not produce an adverse, allergic or other untoward reaction when administered to an animal, such as, for example, a human, as appropriate. The preparation of a pharmaceutical composition that contains at least one herpes virus vector or additional active ingredient will be known to those of skill in the art in light of the

present disclosure, as exemplified by Remington's Pharmaceutical Sciences, 18th Ed., Mack Printing Company, 1990, incorporated herein by reference. Moreover, for animal (*e.g.*, human) administration, it will be understood that preparations should meet sterility, pyrogenicity, general safety and purity standards as required by the U.S. FDA Office of Biological Standards or equivalent governmental regulations in other countries, where applicable.

As used herein, "pharmaceutically acceptable carrier" includes any and all solvents, dispersion media, coatings, surfactants, antioxidants, preservatives (*e.g.*, antibacterial agents, antifungal agents), isotonic agents, absorption delaying agents, salts, preservatives, drugs, drug stabilizers, gels, binders, excipients, disintegration agents, lubricants, sweetening agents, flavoring agents, dyes, and like materials and combinations thereof, as would be known to one of ordinary skill in the art (see, for example, Remington's Pharmaceutical Sciences, 18th Ed., Mack Printing Company, 1990, pp. 1289-1329, incorporated herein by reference). Except insofar as any conventional carrier is incompatible with the active ingredient, its use in the therapeutic or pharmaceutical compositions is contemplated.

A pharmaceutical composition comprising a herpes virus vector and/or additional agent(s) may exploit different types of carriers depending on whether it is to be administered in solid, liquid or aerosol form, and whether it need be sterile for such routes of administration as injection. The pharmaceutical compositions can be administered intravenously, intradermally, intraarterially, intra-graft, intraperitoneally, intralesionally, intracranially, intraarticularly, intraprostatically, intrapleurally, intratracheally, intranasally, intravitreally, intravaginally, intrarectally, topically, intratumorally, intramuscularly, intraperitoneally, subcutaneously, subconjunctivally, intravesicularly, mucosally, intrapericardially, intraumbilically, intraocularly, orally, topically, locally, inhalation (*e.g.*, aerosol inhalation), injection, infusion, continuous infusion, localized perfusion bathing target cells directly (*e.g.*, in an autogenous tissue graft), via a catheter, via lavage, in cremes, in lipid compositions (*e.g.*, liposomes), or by any other method or any combination of the foregoing as would be known to one of ordinary skill in the art (see, for example, Remington's Pharmaceutical Sciences, 18th Ed., Mack Printing Company, 1990).

The actual dosage amount of a pharmaceutical composition of the present invention administered to an animal patient can be determined by physical and physiological factors such as body weight, severity of condition, the type of disease being treated, previous or concurrent therapeutic interventions, idiopathy of the patient and the planned route of administration. The practitioner responsible for administration will, in any event, determine the concentration of active ingredient(s) in a pharmaceutical composition and appropriate dose(s) for the individual subject using routine procedures.

In certain embodiments, pharmaceutical compositions may comprise, for example, at least about 0.1% of an active compound (*e.g.*, a herpes virus vector, a therapeutic agent). In other embodiments, the active compound may comprise between about 2% to about 75% of the weight of the unit, or between about 25% to about 60%, for example, and any range derivable therein. In other non-limiting examples, a dose may also comprise from about 1 microgram/kg body weight, about 5 microgram/kg body weight, about 10 microgram/kg body weight, about 50 microgram/kg body weight, about 100 microgram/kg body weight, about 200 microgram/kg body weight, about 350 microgram/kg body weight, about 500 microgram/kg body weight, about 1 milligram/kg body weight, about 5 milligram/kg body weight, about 10 milligram/kg body weight, about 50 milligram/kg body weight, about 100 milligram/kg body weight, about 200 milligram/kg body weight, about 350 milligram/kg body weight, about 500 milligram/kg body weight, to about 1000 mg/kg body weight or more per administration, and any range derivable therein. In non-limiting examples of a derivable range from the values provided herein, a range of about 5 mg/kg body weight to about 100 mg/kg body weight, about 5 microgram/kg body weight to about 500 milligram/kg body weight, etc., can be administered, based on the values disclosed.

In any case, the pharmaceutical composition may comprise various antioxidants to retard oxidation of one or more component. Additionally, the prevention of the action of microorganisms can be brought about by preservatives such as various antibacterial and antifungal agents, including, but not limited to, parabens (*e.g.*, methylparabens, propylparabens), chlorobutanol, phenol, sorbic acid, thimerosal or combinations thereof.

A herpes virus vector and/or an agent may be formulated into a pharmaceutical composition in a free base, neutral or salt form. Pharmaceutically acceptable salts include the acid addition salts, *e.g.*, those formed with the free amino groups of a proteinaceous composition, or which are formed with inorganic acids such as for example, hydrochloric or phosphoric acids, or such organic acids as acetic, oxalic, tartaric or mandelic acid. Salts formed with the free carboxyl groups can also be derived from inorganic bases such as sodium, potassium, ammonium, calcium or ferric hydroxides; or such organic bases as isopropylamine, trimethylamine, histidine or procaine.

In embodiments where the pharmaceutical composition is in a liquid form, a carrier can be a solvent or dispersion medium including, but not limited to, water, ethanol, polyol (*e.g.*, glycerol, propylene glycol, liquid polyethylene glycol, etc.), lipids (*e.g.*, triglycerides, vegetable oils, liposomes) and combinations thereof. The proper fluidity can be maintained, for example, by the use of a coating, such as lecithin; by the maintenance of the required particle size by dispersion in carriers such as, for example liquid, polyol or lipids; by the use of surfactants such as, for example, hydroxypropylcellulose; or combinations thereof. In many cases, it will be preferable to include isotonic agents, such as sugars, sodium chloride or combinations thereof.

In other embodiments, one may use eye drops, nasal solutions or sprays, aerosols or inhalants in the present invention. Such compositions are generally designed to be compatible with the target tissue type. In a non-limiting example, nasal solutions are usually aqueous solutions designed to be administered to the nasal passages in drops or sprays. Nasal solutions are prepared so that they are similar in many respects to nasal secretions, so that normal ciliary action is maintained. Thus, in preferred embodiments, the aqueous nasal solutions usually are isotonic or slightly buffered to maintain a pH of about 5.5 to about 6.5. In addition, antimicrobial preservatives, similar to those used in ophthalmic preparations, drugs, or appropriate drug stabilizers, if required, may be included in the formulation. For example, various commercial nasal preparations are known to include drugs such as antibiotics or antihistamines.

In certain embodiments, the viral vector concentration that may be administered is about 10^2 , about 10^3 , about 10^4 , about 10^5 , about 10^6 , about 10^7 ,

about 10^8 , about 10^9 , to about 10^{10} PFU/ml, and any integer derivable therein, and any range derivable therein. In some embodiments, the concentration range of viral vectors that may be administered to a tissue or organism may include from about 10^3 , about 10^4 , about 10^5 , about 10^6 , about 10^7 , about 10^8 , about 10^9 , about 10^{10} , about 10^{11} , about 10^{12} , to about 10^{13} PFU/kg, and any integer derivable therein, and any concentration range derivable therein. In other embodiments, the concentration range of viral vectors that may be administered to a cell (MOI) may include from about 0.1, about 0.2, about 0.3, about 0.4, about 0.5, about 0.6, about 0.7, about 0.8, about 0.9, about 1, about 2, about 3, about 4, about 5, about 6, about 7 about 8, about 9, about 10, about 11, about 12, about 13, about 14, about 15, about 16, about 17, about 18, about 19, about 20, about 21, about 22, about 22, about 23, about 24, about 25, about 26, about 27, about 28, about 29, about 30, about 31, about 32, about 33, about 34, about 35, about 36, about 37, about 38, about 39, about 40, about 41, about 42, about 43, about 44, about 45, about 46, about 47, about 48, about 49, about 50, about 51, about 52, about 53, about 54, about 55, about 56, about 57, about 58, about 59, about 60, about 61, about 62, about 63, about 64, about 65, about 66, about 67, about 68, about 69, about 70, about 71, about 72, about 73, about 74, about 75, about 76, about 77, about 78, about 79, about 80, about 81, about 82, about 83, about 84, about 85, about 86, about 87, about 88, about 89, about 90, about 91, about 92, about 93, about 94, about 95, about 96, about 97, about 98, about 99, to about 100 PFU/cell, and any range derivable therein.

In certain embodiments, the pharmaceutical composition is prepared for administration by such routes as oral ingestion. In these embodiments, the solid composition may comprise, for example, solutions, suspensions, emulsions, tablets, pills, capsules (*e.g.*, hard- or soft-shelled gelatin capsules), sustained release formulations, buccal compositions, troches, elixirs, suspensions, syrups, wafers, or combinations thereof. Oral compositions may be incorporated directly into the food of the diet. Preferred carriers for oral administration comprise inert diluents, assimilable edible carriers or combinations thereof. In other aspects of the invention, the oral composition may be prepared as a syrup or elixir. A syrup or elixir may comprise, for example, at least one active agent and, optionally, a

sweetening agent, a preservative, a flavoring agent, a dye, a preservative, or combinations thereof.

In certain embodiments, an oral composition may comprise one or more binders, excipients, disintegration agents, lubricants, flavoring agents, and combinations thereof. The composition may comprise one or more of the following: a binder, such as gum tragacanth, acacia, cornstarch, gelatin or combinations thereof; an excipient, such as dicalcium phosphate, mannitol, lactose, starch, magnesium stearate, cellulose, magnesium carbonate or combinations thereof; a disintegrating agent, such as corn starch, potato starch, alginic acid or combinations thereof; a lubricant, such as magnesium stearate; a sweetening agent, such as sucrose, lactose, saccharin or combinations thereof; a flavoring agent, such as peppermint, oil of wintergreen, cherry flavoring, orange flavoring, and the like; or combinations thereof. When the dosage unit form is a capsule it may contain, in addition to materials of the above type, carriers such as a liquid carrier. Various other materials may be present as coatings or to otherwise modify the physical form of the dosage unit. For instance, tablets, pills, or capsules may be coated with shellac, sugar or both.

Additional formulations which are suitable for other modes of administration include suppositories. Suppositories are solid-dosage forms of various weights and shapes, usually medicated, for insertion into the rectum, vagina or urethra. After insertion, suppositories soften, melt or dissolve in the cavity fluids. In general, for suppositories, traditional carriers may include, for example, polyalkylene glycols, triglycerides or combinations thereof. In certain embodiments, suppositories may be formed from mixtures containing, for example, the active ingredient in the range of about 0.5% to about 10%, and preferably about 1% to about 2%.

Sterile injectable solutions are prepared by incorporating the active compounds in the required amount in an appropriate solvent with various other ingredients as disclosed above, as required, followed by filter sterilization. Generally, dispersions are prepared by incorporating the various sterilized active ingredients into a sterile vehicle which contains the basic dispersion medium and/or the other ingredients. In the case of sterile powders for the preparation of sterile injectable solutions, suspensions or emulsions, the preferred methods of preparation are vacuum-drying or freeze-drying techniques which yield a powder of the active

ingredient, optionally with any additional desired ingredient in solid or liquid form. The liquid medium may be suitably buffered, as appropriate, and the liquid composition is rendered isotonic with sufficient saline or glucose where injection is the mode of administration. The preparation of highly concentrated compositions for direct injection is also contemplated, where the use of DMSO as solvent is envisioned to result in extremely rapid penetration, delivering high concentrations of the active agents to a small area.

The composition is preferably stable under the conditions of manufacture and storage, and is preferably preserved against the contaminating action of microorganisms, such as bacteria and fungi. It will be appreciated that endotoxin contamination is desirably kept minimally at a safe level, for example, less than 0.5 ng/mg protein.

In particular embodiments, prolonged absorption of an injectable composition can be brought about by the use in the compositions of agents delaying absorption, such as aluminum monostearate, gelatin or combinations thereof.

G. KITS

Any of the compositions described herein may be formulated into a kit. In a non-limiting example, a vector (*e.g.*, a herpes virus vector comprising a therapeutic nucleic acid for a vascular or cardiovascular disease or disorder) and optionally an additional agent, may be formulated into a kit. The kits will thus contain, in suitable container means, a vector and, optionally, an additional agent.

The kits may comprise a suitably aliquoted vector and optional additional agent composition, whether labeled or unlabeled, which may be used, *e.g.*, to prepare a standard curve for a detection assay. The components of the kits may be packaged either in aqueous media or in solid form, *e.g.*, lyophilized form. The container means of the kits will generally include at least one vial, test tube, flask, bottle, syringe or other container means, into which a component may be placed, and preferably, suitably aliquoted. Where there is more than one component in the kit, the kit also will generally contain a second, third or other additional container into which the additional components may be separately placed. However, various combinations of components may be contained in a vial. The kits of the present invention also will typically include a means for containing the vector and optional

additional agent, and any other reagent containers, in close confinement for commercial sale. Such containers may include injection- or blow-molded plastic containers into which the desired vials are retained.

5 When the components of the kit are provided in one or more liquid solutions, the liquid solution is an aqueous solution, with a sterile aqueous solution being particularly preferred. The vector and optional additional agent composition may also be formulated into a syringeable composition. In the case of syringeable compositions, the container means may itself be a syringe, pipette, and/or other similar apparatus, from which the formulation may be applied to an area of the
10 body, injected into an animal, or applied to, and/or mixed with, the other components of the kit.

The components of the kit also may be provided as dried powder(s). When reagents and/or components are provided as a dry powder, the powder can be reconstituted by the addition of a suitable solvent or liquid carrier in this case of
15 mixtures. It is envisioned that the solvent may also be provided in another container means.

Irrespective of the number and type of containers, the kits of the invention may also comprise, and be packaged with, an instrument for assisting with the injection/administration of a pharmaceutical composition of the present invention
20 within a tissue, organ or body of an animal. Such an instrument may be a syringe, pipette, forceps, or any medically approved delivery device.

The following examples are included to demonstrate preferred embodiments of the invention. However, those of skill in the art will, in light of the present disclosure, appreciate that many alterations can be made in the specific embodiments
25 which are disclosed herein without departing from the spirit and scope of the invention as claimed.

EXAMPLE 1

VECTOR PREPARATION

30 R3616 and R849

Described herein is the construction of a replication competent, neurovirulence-attenuated (debilitated for growth within the central nervous system), recombinant γ 134.5-deleted HSV-1 virus containing the lacZ gene under the control

of the γ 134.5 promoter (HSVlacZ or R849). The following methods and vectors are set forth in U.S. Patent Nos. 5,238,688 and 6,120,773, both incorporated by reference in their entireties.

Although the HSV vector R3616 exemplifies a growth-debilitated HSV via disruption in a non-essential gene or genes, those of skill in the art will recognize that other non-essential HSV genes may be disrupted (*e.g.*, insertion, substitution or deletion) to obtain a similar growth-debilitated HSV (see Table 1 and 3). Further, those of skill will also recognize that such a disrupted gene may be an essential gene (see Tables 1 and 2).

A recombinant virus R3617, lacking 1 Kbp of DNA in each copy of the γ 134.5 gene, was generated by co-introduction into rabbit skin cells of intact R4002 DNA and the DNA of plasmid pRB3616. In plasmid pRB3616, the sequences containing most of the coding region of a γ 134.5 gene, *i.e.*, those located between the BstEII and StuI sites within the BamHI S fragment of HSV-1 strain F, had been deleted. To generate pRB3616, plasmid pRB143 [Post *et al.*, *Proc. Nat'l. Acad. Sci. (USA)*, 7.,7, 4201 (1980)] was digested with BstEII and StuI, blunt-ended with T4 polymerase, and religated. The progeny of the transformation were plated on 143 *tk*⁻ (thymidine kinase-) cells overlaid with medium containing BUdR (bromodeoxyuridine) to select for *tk* viruses. Because the *tk* gene is present in both copies of the γ 134.5 gene in R4002, the selected progeny contain deletions in both copies. The selected *tk*⁻ virus, designated R3617, was analyzed for the presence of the deletion in both copies of the γ 134.5 gene. An intact γ 134.5 gene was not present.

For assays of neurovirulence, the deletion in the native *tk*⁻ gene of R3617, which traces its origin from HSV-1(F) Δ 305, had to be repaired. This was accomplished by co-introduction into rabbit skin cells of intact R3617 DNA and Bam HI Q fragment containing the *tk* gene (Chou and Roizman, 1994). The virus selected for a *tk*⁺ phenotype in 143*tk*⁻ cells was designated R3616 (Chou *et al.*, 1990). The recombinant R849 (HSVlacZ) virus carrying the β -galactosidase reporter cassette in place of the γ 134.5 genes was generated as follows. Rabbit skin cells were transfected with intact R4002 DNA and a plasmid having the BstEII-StuI DNA fragment of the γ 134.5 replaced with a BstEII-StuI fragment containing the coding domain of the β -galactosidase gene. Recombination occurred through the

flanking sequences. The progeny of the transfection was selected in 143tk cells in the presence of BUdR. A clone was purified, tested for the presence of the β -galactosidase open reading frame, and then contrransfected with BamHI Q DNA fragments to rescue the thymidine kinase gene. In the R849 (HSVlacZ) recombinant virus, the β -galactosidase coding sequence is fused in frame with the γ 134.5 gene coding for the first 28 amino acids of the gene product. The chimeric gene is expressed from the γ 134.5 promoter. Viral titers were measured in Vero cells. R3616 was deposited under accession number ATCC VR2280 with the American Type Culture Collection, Manassas, Virginia on Aug. 14, 1990.

Adeno-associated Viral Vectors

For controls in comparative experiments, adeno-associated viral (AAV) vectors were created and purified as previously described (Svensson et al., 1999). Briefly, the AAV helper plasmid, pAd14 contains the AAV rep and ap genes cloned into the Xba I site, and the SV40 origin of replication cloned into the BamHI site of pBluescript. The virus was purified by CsCl gradient centrifugation, dialyzed against PBS, and stored in aliquots at -80°C. Viral titer was determined by using a dot blot hybridization assay to determine the number of viral genomes/ml. HeLa cells were infected with the virus and detected by staining with 5-bromo-4-chloro-3-indolyl-beta-D-galactopyranoside (X gal) to reveal β -galactosidase (lacZ) activity 24 hours after infection.

EXAMPLE 2

CELL CULTURE AND MTT ASSAY

To determine the appropriate dose for vascular gene transfer, human umbilical vein endothelial cells (HUVECs) were contacted with graded concentrations of HSV-1 for 120 min and viability was assessed using a standard 72 hr MTT assay, as described below.

Human umbilical vein endothelial cells (HUVECs; BioWhittaker/Clonetics) were suspended in complete medium EBM-2 (BioWhittaker #CC3156) and singlequots of EGM-2 (#CC4176) and grown to 100% confluence in P75 flasks in a 37°C incubator with humidified 5% CO₂. For the passage of cells, all reagent pack contents (HBSS, Trypsin/EDTA, TNS; BioWhittaker CC-5034) were thawed in a 37°C water bath and exhausted media was removed and discarded. HUVECs were

washed with enough HBSS to cover them, trypsin/EDTA was added, and flasks were gently agitated under an inverted microscope. Following loss of adherence of cells, trypsin was inactivated with an equal volume of TNS. The cells were centrifuged at 1200 rpm for 5 min and the pellet was resuspended in complete EBM-2 medium and singlequots of EGM-2.

A modified colorimetric assay based on the selective ability of living cells to reduce the yellow salt MTT (3-[4,5-dimethylthiazol-2-yl]-2,5 diphenyl tetrazolium bromide; Sigma Chemical Co.) to formazan was used to quantitate HUVEC viability following viral exposure. HUVECs were suspended in media and plated into 96-well flat-bottom microtiter plates at a density of 5×10^3 per ml in a volume of 0.1 ml. Cells were allowed to incubate overnight and then exposed to HSV R3616 suspended in HEPES-buffered saline (HBS) for 2 hr at 37°C with humidified 5% CO₂. The supernatants were aspirated, and 0.1 ml of complete medium EBM-2 was added to each well. The zero plate was allowed to incubate for 60 min. The supernatant was again aspirated and MTT (5 mg/ml) was added to each well of the zero plate for 4 hr at 37°C. After formation of formazan crystals, the supernatant was aspirated with attention not to disturb the underlying cell layer. The formazan crystals were then dissolved in 0.1 ml DMSO (Sigma) and the absorbance measured at 515 nm using a microplate spectrophotometer (SpectraMax 340 004) interfaced with a personal computer (Apple Macintosh). This process was repeated after 72 hours and relative cell survival was calculated. The experiment was repeated a total of three times.

HUVECs infected with R3616 at an MOI = 1 survived over 72 hr at a rate of 92±19%. As expected, a decreasing proportion of cells survived with increasing viral concentrations, with only 4±2% cells surviving at an MOI = 50. Subsequent viral doses were chosen after approximation of cell number in whole-tissue volumes with MOIs ranging from 1-2.

EXAMPLE 3

INFECTION OF HUMAN SAPHENOUS VEINS

IN WHOLE ORGAN CULTURE

To test *in vitro* gene transfer in quiescent vascular tissue, human saphenous veins discarded at the time of operation were infected at 120 mmHg for 10 min with

vehicle, HSVlacZ (R849), or an adeno-associated virus control containing the lacZ gene (AAVlacZ).

Segments of saphenous vein were harvested from patients at the time of coronary or peripheral bypass surgery under an approved protocol. No patient had any history of venous disease. Human saphenous vein segments were judged to be of good quality by the operating surgeon with diameters ranging from 3.5 to 5.5 mm. Human saphenous veins were infected and cultured whole.

Segments were gently irrigated with saline to remove any residual debris, and excess adventitia trimmed sharply. Vein segments were cut into rings and placed in separate culture wells with RPMI-1640 media/15% FBS and incubated at 37°C. The vein was divided into adjacent control and experimental segments and separately dually cannulated with 22 gauge catheters at both ends for vector infusion and pressure monitoring (Hewlett-Packard 78534B; Palo Alto, CA). Veins were distended with either vehicle (n=4), AAVlacZ 4×10^{11} pfu/ml (n=2), or HSVlacZ (R849) 6.5×10^8 pfu/ml for 10 min at 120 mmHg with limited outflow and low shear stress. The main channel of the vein was then rinsed with saline and the vein cut into rings 0.5 cm in length.

The first ring was fixed immediately in 1.25% glutaraldehyde for 10 min, washed in PBS and incubated overnight at 37°C with X-gal in a dark environment for chromogenic assay. The remaining rings (4-7, depending on the amount of vein available) were each placed in separate culture wells with six ml media (RPMI-1640 with 15% FBS) and incubated at 37° C. Media was changed every two days. Vein segments were removed from the media, fixed, and stained with X gal as above after 3, 5, 10, 14, 20, 28, 32, 34, 40, 48, and 55 days.

All segments exposed to HSVlacZ (R849) expressed β -galactosidase. Expression was not present immediately after exposure in any veins, but after 20 days (*e.g.*, 34 days) all veins showed significant expression in the intima and adventitia and, to a lesser extent, the media. Segments were counterstained with eosin. HSV-mediated transgene expression was not limited to the EC layer, but was present in all cell layers, including a significant amount of expression in the medial layer, which appeared to exceed that observed with adenoviral-mediated infection. The ability of HSV to effectively penetrate the EC layer and infect cells deep within the vein wall was also surprising, as it had not been previously demonstrated.

Expression persisted to 55 days, the final time point in this study. This is in stark contrast to adenoviral-mediated gene transfer, which generally dissipates after seven days.

Expression after AAVlacZ exposure was minimal, with saphenous veins exposed to AAVlacZ exhibiting inconsistent staining in the intima and adventitia, which was much less pronounced than with HSVlacZ (R849). Expression was not evident in any vein segment exposed to vehicle only.

Sections were also stained with a rabbit anti-human antibody to von Willebrand Factor (vWF) to examine the presence and integrity of the endothelium by immunohistochemistry. 4 μ m sections of paraffin-embedded tissues were mounted on poly-L-lysine coated slides, and deparaffinized by sequential 5 min washes with xylene and decreasing concentrations of ethanol and dH₂O. Sections were then incubated in pre-warmed 10mM EDTA (pH 6.0) for 60 min at 70°C, washed three times with dH₂O, and incubated in pre-warmed Proteinase K (20 μ g/ml in PBS) for 20 min at 37°C. After blocking with a 10% solution of normal goat serum (Vector, Burlingame, CA) in 0.5% casein (Sigma) and PBS, endogenous avidin and biotin were quenched using a blocking kit (Vector). Slides were washed three times with 0.5% casein/0.01% Tween 20 in PBS and incubated overnight at 4°C with rabbit anti-human vWF (DAKO, Denmark) titered to 0.8 μ g/ml in 0.5% casein. Appropriate negative controls (rabbit IgG isotype, DAKO) were also used. Slides were washed three times and incubated with goat-anti-rabbit IgG biotinylated secondary antibody (2 μ g/ml in PBS, 0.5% casein and 10 % normal human serum) for 25 min at 37°C. Endogenous peroxidase was quenched with 3% H₂O₂ (20 min at room temperature (RT)), and slides were washed and incubated with Vectastain Standard Elite (ABC) for 25 minutes at RT. Tissues were developed with the Vector DAB kit, and counterstained with a 0.03% solution of light green SF yellowish (Fisher, Hanover Park, IL). Sections were dehydrated in alcohol and xylene, mounted with Permount and coverslips, and examined via light microscopy.

Treatment with HSVlacZ (R849) did not appreciably affect the integrity of the EC layer as assessed qualitatively by vWF staining in cultured human veins 34 days after exposure to vehicle or HSVlacZ (R849).

EXAMPLE 4

INFECTION OF RABBIT EXTERNAL JUGULAR VEIN PATCHES

To demonstrate *in vivo* gene transfer in proliferating vascular tissue, the external jugular vein of male New Zealand White rabbits was exposed to vehicle, HSVlacZ (R849), or AAVlacZ.

Male New Zealand white rabbits (3.5-4 kg) were anesthetized by intramuscular (1m) injection with ketamine hydrochloride (40 mg/kg) and xylazine (5 mg/kg) augmented with halothane via endotracheal intubation. Antibiotic prophylaxis was provided with enrofloxacin 10 mg/kg IM daily. Using sterile technique, the external jugular vein was exposed and two branches cannulated with 24-gauge catheters. One cannula was used for irrigation and infection, and the other for intraluminal pressure monitoring. The main channel was infected with either vehicle (n=6), AAVlacZ 4×10^{11} pfu/ml (n=2), or HSVlacZ (R849) 4×10^8 pfu/ml (n=4) for 10 min at 100 mmHg.

Following infection, the vein was irrigated with saline, excised and bivalved. The ipsilateral common carotid artery (CCA) was then exposed through the same incision and the animal systemically anticoagulated with heparin (200 U/kg) intravenously. The CCA was doubly clamped and a 1.5-cm longitudinal arteriotomy made proximal to the cranial thyroid branch. The arteriotomy was reconstructed with external jugular vein patch angioplasty using running 8-0 polypropylene suture (Davis-Geck TE-145; Manatee, PR) under 4.5x-loupe magnification. Ultrasonic transit-time flow through the graft was measured (Transonics Systems Inc. HT207; Ithaca, NY) and recorded using a digital data acquisition system (Lab Master D.M.A.; Scientific Solutions, Inc., Solon, OH). All operations for implantation of external jugular vein patches in rabbits were completed without complication. There was no significant difference in mean blood flow in vehicle-treated vs. viral-infected patches at time of implantation (vehicle 15.0 ± 1.2 vs. viral 18 ± 1.0 ml/min; p=NS). The incision was closed and the animal allowed to recover. Anticoagulation was not reversed.

After four weeks, the vein patches were harvested and assessed for patency and β -galactosidase expression using X gal. The animals were reanesthetized and the vein patches reexposed. Intraarterial pressure and blood flow through the patch

were again measured and recorded. All vein patches were patent at the time of harvest at four weeks (n=12).

The vein patch was divided into three sections starting one mm distal to the beginning of the patch and ending one mm proximal to its endpoint. The harvested vessels were fixed in 1.25% glutaraldehyde for 10 min, washed in PBS, and incubated overnight at 37°C with X gal. Five μ m sections were cut from each of the portions of the patch and stained using eosin, hematoxylin and eosin (H & E) and the Weigert van Gieson. Ten high-power fields (40X) of each tissue section were examined using a microscope (Nikon Microphoto-FX) equipped with a digital camera (Sony PowerHAD 3CCD). Stained cells/nuclei were counted using standard imaging software (ImagePro Plus 3.0.1). The total number of β -galactosidase positive cells was counted in the neointima, media and adventitia located circumferentially about the lumen. The total number of cells was counted on the H & E sections in the neointima, media and adventitia. The average number of positive cells was expressed as a percentage of the total number of cells, reflecting the infection efficiency.

All vein patches that had been exposed to HSVlacZ (R849; n=4) showed significant β -galactosidase expression in all layers of the vein wall at 4 weeks after exposure (*i.e.*, infection), especially within the smooth muscle cells (SMCs) comprising the neointima ($48 \pm 2\%$ infection efficiency). All slides were counterstained with eosin. The calculated infection efficiencies in the adventitia, media, and neointima were $42 \pm 1\%$, $58 \pm 3\%$ and $48 \pm 2\%$, respectively. Interestingly, there was also moderate expression in the arterial segment (the posterior wall of the reconstruction), especially near the margins surrounding the suture artifact. In contrast, patches infected with AAVlacZ showed inconsistent transgene expression, mostly confined to the adventitia. Expression was not evident in the vehicle-exposed patches or in any harvested external jugular veins or CCAs contralateral to an HSVlacZ (R849)-infected vein patch or an AAV-infected vein patch.

EXAMPLE 5

EFFECTS OF R849 ON CULTURED HUMAN CELLS

To demonstrate the effect of HSV R849 (HSVlacZ) in cultured human cells, human umbilical vein endothelial cells (HUVECs; BioWhittaker/Clonetics

#CC2517) and umbilical artery smooth muscle cells (#CC2579) were separately suspended in complete medium EGM bullet kit (#CC3182) and grown to 100% confluence in P75 flasks in a 37°C incubator with humidified 5% CO₂. A modified colorimetric assay based on the selective ability of living cells to reduce the yellow salt MTT (3-[4,5-dimethylthiazol-2-yl]-2,5 diphenyl tetrazolium bromide; Sigma Chemical Co.) to formazan was used to quantitate HUVEC and UASMC viability following viral exposure and rescue with Acyclovir (Abbott Laboratories).

HUVECs infected with R849 (HSVlacZ) at an MOI=1 had relative cell survivals of 102±10% and 81±4% cells after 48 and 72 hr, respectively. Similarly infected UASMCs had relative cell survivals of 92±10% and 73±6%, respectively. As expected, a decreasing proportion of cells survived with increasing viral concentrations, with only 12±3% of HUVECs and 5±10% of UASMCs surviving after infection with an MOI=25 (relative 72 hr cell survival with Acyclovir (0.25 µg/ml) was 45±5%; with Acyclovir (0.5 µg/ml) was 85±9%) while similarly infected UASMCs could not be rescued, even with 10 µg/ml Acyclovir. These results demonstrate that (1) R849 (HSVlacZ) is relatively non-toxic at low doses, (2) R849 (HSVlacZ) is more toxic to SMCs than endothelial cells (ECs) at higher doses, and (3) a majority of ECs exposed to an MOI=25 can be rescued with therapeutic doses of Acyclovir, while SMCs cannot. Thus, R849 (HSVlacZ) exhibits selectivity for SMCs, the target cell population in this therapeutic strategy.

EXAMPLE 6

EFFECTS OF R849 ON WHOLE HUMAN VEINS

Effects of R849 (HSVlacZ) on whole human veins were again studied (as described above in Example 3). Unused portions of saphenous veins harvested at the time of coronary or peripheral bypass were cannulated with 22 gauge catheters at both ends and distended for 10 min at 120 mmHg with either vehicle or 1x10⁹ PFU/ml R849 (HSVlacZ; n=8). The first ring was fixed immediately in 1.25% glutaraldehyde for 10 min, washed in PBS, and incubated overnight at 37°C with X gal in a dark environment. The remaining rings were each placed in separate culture wells with 6 ml media (RPMI-1640 with 30% FBS) and incubated at 37°C. Vein segments were removed from media, fixed, and stained with X gal after 2-55 days.

5 The results showed that R849 (HSVpacZ)-infected segments exhibited β -galactosidase expression up to 55 days following infection. Interestingly, transgene expression was not limited to the ECs but was present in all cell layers, including a significant amount of expression in the media, which appeared to exceed that observed with adenoviral-mediated infection.

10 Obviously, any strategy designed to limit restenosis must not disturb the antithrombotic endothelial layer. To test for possible negative effects of HSV on the endothelium, cultured human saphenous veins were exposed to R849 (HSVlacZ) and then examined for the endothelial marker, von Willebrand factor (vWF), by immunostaining. Veins were distended with either vehicle or R849 (HSVlacZ; 6.5×10^8 pfu/ml) for 10 min at 120 mmHg, as described above. Vein segments were removed from the media, fixed, and stained. Four μ m sections of paraffin-embedded tissues were mounted on poly-L-lysine coated slides, and deparaffinized by sequential 5 min washes with xylene and decreasing concentrations of ethanol and dH₂O. Slides were washed three times with 0.5% casein/0.01% Tween 20 in PBS and incubated overnight at 4°C with rabbit anti-human vWF (DAKO, Denmark) titrated to 0.8 μ g/ml in 0.5% casein. Appropriate negative controls (rabbit IgG isotype, DAKO) were also used. Slides were washed three times and incubated with goat anti-rabbit IgG biotinylated secondary antibody (2 μ g/ml in PBS, 0.5% casein and 10% normal human serum) for 25 min at 37°C. Tissues were developed with the Vector DA kit, and counterstained with a 0.03% solution of light green SF yellowish (Fisher, Hanover Park, IL). Gene transfer was consistently demonstrated throughout the vascular wall, with the endothelium remaining largely intact as judged by the presence of vWF.

25 EXAMPLE 7

R849 INFECTION IN VEIN GRAFTS IN IMMUNOCOMPETENT ANIMALS

30 Experiments in the rabbit vein patch model were again performed (as described in Example 4) in order to examine the time course of R849 (HSVlacZ)-mediated infection, with the potential for long-term programmed expression.

Male New Zealand White rabbits (3.5-4 kg) were anesthetized and the external jugular vein was surgically exposed. The main channel was infected with

either vehicle (n=6), R849 (HSVlacZ) 4×10^8 pfu/ml (n=8) or R3616 3×10^8 pfu/ml
 (control virus, n=2) for 10 min at 120 mmHg. Following infection, the vein was
 excised and fashioned as a common carotid vein patch angioplasty using running 8-0
 polypropylene suture. This vein patch technique has the advantage of creating an
 5 arterialized vein segment with minimal surgical trauma. After four weeks, the
 animals were reanesthetized and the vein patches re-exposed. There was no
 significant difference in mean blood flow in control (31 ± 2 ml/min) vs. viral-
 infected patches (34 ± 3 ml/min) at the time of implantation. All vein patches were
 patent at the time of harvest at four weeks. Patches exposed to R849 (HSVlacZ)
 10 4×10^8 pfu/ml showed marked β -galactosidase staining in all parts of the vessel wall,
 most notably within the medial layer. Calculated infection efficiencies (percent of
 cells infected) were $42 \pm 2\%$ in the adventitia, $44 \pm 9\%$ in the media, and $45 \pm 6\%$ in
 the neointima. Moderate β -galactosidase staining was also detected in the untreated
 arterial segment adjacent to the vein patch, and in the more distal portions of the
 15 native carotid artery. These consistent rates of infection, coupled to stable and
 durable transgene expression, far exceed the reported results with other
 vectors.[Moawad *et al.*, *Ann. Vasc. Surg.* **15**:367-73, 2001; Schwartz *et al.*, *J.*
Vasc. Surg. **29**:874-83, 1999; Hanna *et al.*, *J. Vasc. Surg.* **31**:770-80, 2000;
 George *et al.*, *Circ.* **101**:296-304, 2000; Eslaim *et al.*, *J. Vasc. Surg.* **31**:1149-59,
 2000; Dollery *et al.*, *Circ.* **99**:3199-205, 1999; Scheinman *et al.*, *J. Vasc. Surg.*
 20 **29**:360-9, 1999; Schneider *et al.*, *J. Vasc. Surg.* **29**:543-50, 1999, and Waugh *et*
al., *Circ.* **102**:332-7, 2000]. Furthermore, the dose of R849 (HSVlacZ) required to
 achieve these results, no greater than
 10⁸ pfu/ml, is several orders of magnitude less than traditional strategies using
 25 adenovirus (see above).

EXAMPLE 8

EFFECT OF R849 ON NEOINTIMAL HYPERPLASIA IN IMMUNOCOMPETENT ANIMALS

30 A novel model of hemodynamically compromised vein patch angioplasty was
 used to test the effects of R849 (HSVlacZ) on SMC proliferation and neointimal
 hyperplasia. Compromise of blood flow, with its attendant reductions in shear
 stress, is a well-known stimulator of SMC proliferation [Rectenwald *et al.*, *Circ.*

102, 2000 and Kumar *et al.*, *Arterioscler Thromb. Vasc. Biol.* 17:2238-44, 1997], and is an important predisposing factor to vein graft failure [Meyerson *et al.* *J. Vasc. Surg.* 34:90-7, 2001]. Therefore, a model of venous transplantation under profoundly reduced flow was developed in order to simulate failing bypass grafts.

5 Generally, rabbits underwent surgical vein exposure, R849 (HSVlacZ) infection, and vein patch angioplasty as described in the preceding examples. Prior to wound closure, the main branch of the angioplasty reconstruction (the common carotid artery) was ligated, leaving the only outflow via a small cranial thyroid artery. This had the reproducible effect of decreasing blood flow to approximately
10 5% of baseline values, a condition that persisted to sacrifice.

The resultant vein patch was grossly thickened and contracted, the thickest ever reported for an animal model of vein grafting. The histological characteristics were similar to those of failing human bypass grafts, including the presence of laminar thrombus and flow-limiting luminal stenosis.

15 Specifically, this model was used to test the effects of R849 (HSVlacZ) on SMC proliferation *in vivo*. Rabbit external jugular veins were infected with either R849 (HSVlacZ; n=8) or vehicle (PBSS; n=6) for 10 min and then fashioned as carotid vein patches with limited outflow. Mean blood flow and shear stress at the time of implantation were similar in R849 (HSVlacZ)-infected (2.8 ± 0.3 ml/min, 0.17 ± 0.03 dyne/cm²) and vehicle (2.6 ± 1.7 ml/min, 0.32 ± 0.1 dyne/cm²) groups.
20 After four weeks, the vein patches were re-exposed, perfusion-fixed at the last recorded blood pressure, and morphometrically examined for neointimal hyperplasia. Five μ m sections were cut, stained using the Weigert van Gieson technique to highlight the internal elastic lamina, and analyzed using digital planimetry (Adobe Photoshop; Adobe Systems, Inc.; San Jose, CA). R849
25 (HSVlacZ)-infected patches were widely patent, with significantly less neointimal hyperplasia than vehicle-treated controls (R849 (HSVlacZ) 280 ± 28 vs. vehicle 410 ± 38 μ m, $p=0.003$). Treatment with R849 (HSVlacZ) essentially returned the "low-flow" patch morphology to "high-flow" physiology.

30

EXAMPLE 9

ERADICATION OF ESTABLISHED R849 INFECTION WITH ACYCLOVIR AND SYSTEMIC TOXICITY OF R849 INFECTION OF R849

5 The ability to attenuate or eliminate infection after completion of the desired biological effect of the vector would be a significant enhancement to gene therapy strategies, and would serve to limit toxicity and the potential for long-term sequelae. Acyclovir sodium (12-amino-1,9-dihydro-9-[(2-hydroxy-ethoxy)methyl]-6H-purin-6-one; trade name *Zovirax*; Alphaparm USPD, Inc.) is a synthetic purine nucleoside analog with an antiviral spectrum limited to herpes viruses including herpes simplex, herpes zoster (shingles), varicella (chicken pox), Epstein-Barr virus, and cytomegalovirus. Clinically, its use is limited to the treatment of herpes simplex, genitalis, and zoster infections.

15 Its mechanism of action is believed to include penetration intracellularly, conversion initially to the monophosphate by viral thymidine kinases, then to the diphosphate by cellular guanylate kinase and, finally, to the triphosphate by various other cellular enzymes. Fully active Acyclovir triphosphate competes with the natural substrate, deoxyguanosine triphosphate, for a position in the DNA chain of the herpes virus. Once incorporated, it terminates DNA synthesis via (1) competitive inhibition of viral DNA polymerase, (2) incorporation into, and termination of, the growing viral DNA chain, and (3) inactivation of the viral DNA polymerase. Uninfected cells show only minimal uptake and phosphorylation of Acyclovir. The HSV strains used in this example retain *tk* activity and are thus highly susceptible to neutralization by Acyclovir. To test whether an established HSV-1 infection could be eradicated after completion of the desired therapeutic effect (of the recombinant HSV) *in vivo*, a rabbit model was used. Following surgical exposure, the external jugular vein was infected with R849 (HSVlacZ) at 4×10^8 pfu/ml ($n=8$; 0.2-4.0 ml) for 10 min at 120 mmHg and then fashioned as a vein patch atop the common carotid artery as previously described (vein patch angioplasty). Following one week of recovery, half the rabbits ($n=4$) received Acyclovir (Abbott Laboratories) 75 mg/kg body weight intravenously for a total of five days. Starting on the same post-operative day, rabbits were simultaneously treated with oral Acyclovir (Alphaparm USPD Inc.; 3 mg/cc H₂O), which was maintained until sacrifice. Baseline serum Acyclovir, electrolyte, blood urea

nitrogen, and creatinine levels were measured before initiation of Acyclovir treatment, on days 2, 5, and 10 after initiation of treatment, and at sacrifice. During Acyclovir therapy, serum levels of Acyclovir were maintained within the therapeutic range (9 ± 4 $\mu\text{g/ml}$). There was no significant systemic toxicity as measured by serum electrolytes, blood urea nitrogen, and creatinine levels throughout the study. After four weeks, the animals were reanesthetized and the vein patches re-exposed and harvested. Vessels were fixed in 1.25% glutaraldehyde for 10 min, washed in PBS, and incubated overnight at 37°C with X gal. 5 μm sections were cut and counterstained with eosin. Examination of the sections revealed that Acyclovir treatment completely eliminated reporter gene (lacZ) expression, with β -galactosidase activity evident in <1% of cells ($p < 0.55$). The therapeutic antiproliferative effect appears to be preserved in that treated vein patches remained widely patent even after the R849 (HSVlacZ) infection had been eradicated.

There were no significant aberrations in blood counts, platelet function, coagulation, hepatic or renal function with HSV infection (see Table 6). The infection was localized to the doubly clamped vein so systemic exposure was likely minimal.

Although the present invention has been described in terms of preferred embodiments, it is intended that the present invention encompass all modifications and variations that occur to those skilled in the art upon consideration of the disclosure herein, and in particular those embodiments that are within the broadest proper interpretation of the claims and their requirements. All literature cited herein (scientific articles, U.S. patents, foreign patents, and published patent applications) is incorporated by reference.

Table 6

	WBC	HEMOGLOBIN	PLATELETS
HSV-infected			
Day0 (n=3)	4.8, 10.6, 6.5	12.0, 11.1, 13.4	209, 90, 309
Day1 (n=2)	10.6, 9.3	10.2, 9.4	115, 260
Day7 (n=2)	7.4, 12.9	8.7, 10.7	220, 209
Day14 (n=1)	3.5	11.1	313
Vehicle-infected			
Day0 (n=1)	8.7	11.6	41
Day1 (n=1)	9.5	7.3	136
Day7 (n=1)	9.0	9.7	134
Day14 (n=1)	9.4	10	188
	PT	PTT	
HSV-infected			
Day0 (n=3)	8.1, 5.9, 6.9	18.9, 23.4 (one clumped)	
Day1 (n=2)	5.9, 6.4	6.9, 20.2	
Day7 (n=2)	6.4, 5.6	12.9, 12.9	
Day14 (n=1)	6.4	19.7	
Vehicle-infected			
Day0 (n=1)	7.0	21.2	
Day1 (n=1)	6.2	18.6	
Day7 (n=1)	6.0	12.9	
Day14 (n=1)	6.4	19.7	
	Tbili	SGOT	SGPT
HSV-infected			
Day0 (n=4)	0.3, 0.2, 0.2, 0.1	94, 23, 10, 11	42, 14, 41, 3
Day1 (n=2)	0.2, 0.2	72, 104	85, 71
Day7 (n=2)	0.3, 0.2	12	48
Day14 (n=1)	0.0	9	24
Vehicle-infected			
Day0 (n=1)	0.2	11	31
Day1 (n=1)	0.2	95	60
Day7 (n=1)	0.3	14	30
Day14 (n=1)	0.2	11	23
	BUN	CREATININE	
HSV-infected			
Day0 (n=4)	16, 18, 19, 23	1.7, 1.6, 1.3, 1.9	
Day1 (n=2)	13, 30	1.4, 1.7	
Day7 (n=1)	15	1.2	
Day14 (n=1)	24	1.3	
Vehicle-infected			
Day0 (n=1)	13	1.6	
Day1 (n=1)	21	0.0	
Day7 (n=1)	16	1.5	
Day14 (n=1)	15	1.4	

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